

R 1923 IN



PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C07D 209/34, 209/38, 295/067 C07D 491/113, 403/00 A61K 31/495	A1	(11) International Publication Number: WO 93/12085 (43) International Publication Date: 24 June 1993 (24.06.93)
(21) International Application Number: PCT/SE92/00873 (22) International Filing Date: 16 December 1992 (16.12.92)		(81) Designated States: AT, AU, BB, BG, BR, CA, CH, CS, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, NZ, PL, RO, RU, SD, SE, UA, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, SN, TD, TG).
(30) Priority data: 9103752-3 18 December 1991 (18.12.91) SE		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(71) Applicant: AKTIEBOLAGET ASTRA [SE/SE]; S-151 85 Södertälje (SE). (72) Inventors: BOAR, Bernard, Robin ; 25 Meadow Way, Letchworth, Hertfordshire SG6 3JB (GB). CROSS, Alan, John ; 54 Birchwood Road, West Byfleet, Surrey KT14 6DW (GB). (74) Agents: DANIELSSON, Sten et al.; AB Astra, Patent Department, S-151 85 Södertälje (SE).		
(54) Title: ISATIN DERIVATIVES, PROCESSES FOR THE PREPARATION THEREOF AND PHARMACEUTICAL COMPOSITION COMPRISING THE SAME		
(1)		
<p>(57) Abstract</p> <p>The present invention relates to compounds having general formula (1), wherein: X represents one or more substituents independently selected from hydrogen, lower alkyl, aryl, aryloxy, CN, lower alkoxy, halogen, hydroxy, nitro, trifluoromethyl, alkylsulphonamido, NHCOR where R is lower alkyl or aryl, NR₁R₂ where R₁ and R₂ are independently hydrogen or lower alkyl or together form a ring, CO₂R where R is lower alkyl, or cycloalkyl, cycloalkenyl or bicycloalkyl either optionally further substituted by lower alkyl; Y is CO or CR₃R₄ where R₃ and R₄ are independently hydrogen, lower alkyl, lower alkoxy or together form a cyclic acetal; Z is N or CH; stereo and optical isomers and racemates thereof where such isomers exist, as well as pharmaceutically acceptable acid addition salts thereof and solvates thereof; having therapeutic activity, intermediates for their preparation, processes for their preparation, pharmaceutical formulations containing said compounds and medicinal use of said compounds and similar known compounds.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BG	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	IE	Ireland	PT	Portugal
BR	Brazil	IT	Italy	RO	Romania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SK	Slovak Republic
CI	Côte d'Ivoire	LJ	Liechtenstein	SN	Senegal
CM	Cameroon	LK	Sri Lanka	SU	Soviet Union
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	MC	Monaco	TG	Togo
DE	Germany	MG	Madagascar	UA	Ukraine
DK	Denmark	ML	Mali	US	United States of America
ES	Spain	MN	Mongolia	VN	Viet Nam

Isatin derivatives, processes for the preparation thereof and pharmaceutical composition comprising the same.

The present invention relates to novel compounds having therapeutic activity, intermediates for their preparation, processes for their preparation, pharmaceutical formulations containing said compounds and medicinal use of said compounds and similar known compounds.

10

Background of the invention

A major characteristic of Alzheimer's Disease (Senile Dementia, SDAT) is a marked central cholinergic dysfunction. This cholinergic deficit has been reported to correlate with cognitive impairment (P.T. Francis et al, New Engl. J. Med., 1985, 313, 7). Various attempts to increase central cholinergic activity and thereby reverse the cognitive deficits have, to date, met with only limited success.

There is some evidence that use of the alkaloid physostigmine can, in some cases, be marginally beneficial, but the use of this compound in the clinic is compromised by a low therapeutic ratio, a short half-life and poor bioavailability. The cholinesterase inhibitor, 9-amino-1,2,3,4-tetrahydroacridine (THA) has been reported to be of therapeutic value in the treatment of a small group of patients with SDAT (W.K. Summers et al, New Engl. J. Med., 1986, 315, 1241). Further clinical trials of THA have produced some encouraging results but have been hampered by the association of this drug with certain toxic side effects.

35

Other compounds structurally related to either physostigmine or THA have been reported and are the subject of ongoing investigations.

There remains an urgent need for a safe and clinically effective drug for the symptomatic treatment of Alzheimer's Disease and related conditions.

5 A compound structurally similar to the compounds of the present invention, namely 1-[1-(4-benzyl-piperazinyl)-methyl]isatin, is disclosed in Chemical Abstracts 98(3):16650w referring to Boll Chim. Farm., 1982, 121 (5), pp. 221-9. Said compound is said to have
10 pharmacological activity.

Japanese Patent Application No. 138443/86 (Publication No. KOKAI JP 62-294654A2) discloses 1-[2-(4-benzyl-piperazinyl)ethyl]isatin as an intermediate for the synthesis of isatin derivatives which are useful as an agent for treating gastric or duodenal ulcer of mammals including human beings. Said single compound is deleted from the scope of the present invention by a disclosure in claim 1.

20 Furthermore, European Patent Application EP 0 010 398 relates to isatin derivatives useful for treating allergic symptoms. Among all specific compounds disclosed therein is only one falling within the general formula I of the compounds of the present invention, namely 1-[3-(4-(4-chlorobenzyl)-1-piperazinyl)propyl]-isatin. Said single compound is deleted from the scope of the present invention by a disclosure in claim 1 as well.

30

The present invention

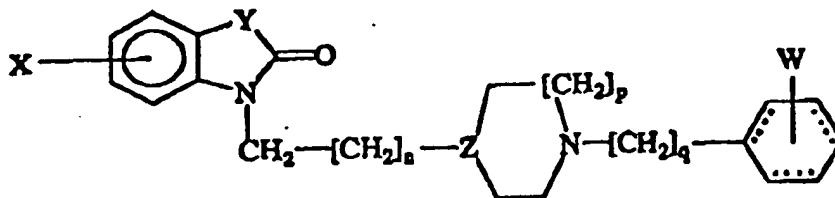
35 A primary objective of the present invention is to provide structurally novel compounds which by virtue of their pharmacological profile enhance cholinergic function and are of value in the treatment of the

cognitiv dysfunctions which may b ass ciated with ageing or with conditions such as Alzheimer's Disease, Senile and related Dementias, Parkinson's Disease, Down's Syndrome and Huntington's Chorea, and in the treatment of conditions such as glaucoma or myasthenia gravis. This utility is manifested, for example, by the ability of these compounds to inhibit the enzyme acetylcholinesterase. Further, the compounds of this invention are, in general, highly potent and selectiv , have an improved duration of action and are, in general, less toxic than hitherto known compounds.

The present invention relates to a compound having the general formula (1)

15

20



(1)

wherein:

25 n is 1, 2 or 3;

p is 1 or 2;

q is 1 or 2;

30

X represents one or more substituents independently selected from hydrogen, lower alkyl, aryl, aryloxy, CN, lower alkoxy, halogen, hydroxy, nitro, trifluoromethyl, alkylsulphonamido,

35

NHCOR where R is lower alkyl or aryl,

NR₁R₂ where R₁ and R₂ are independently hydrogen or lower alkyl or tog ther form a ring,

CO_2R where R is lower alkyl,
or cycloalkyl, cycloalkenyl or bicycloalkyl either
optionally further substituted by lower alkyl;

5 Y is CO or CR_3R_4 where R_3 and R_4 are independently
hydrogen, lower alkyl, lower alkoxy or together form a
cyclic acetal;

Z is N or CH;

10

and  represents an optionally substituted
phenyl or cyclohexyl group; wherein

15

W represents one or more substituents independently
selected from hydrogen, lower alkyl, lower alkoxy or
halogen;

20

stereo and optical isomers and racemates thereof where
such isomers exist, as well as pharmaceutically
acceptable acid addition salts thereof and solvates
thereof;

25

with the provisos that the compound wherein n=1, p=1,

q=1, X=H, Y=CO, Z=N and



= unsubstituted

30

phenyl and the compound wherein n=2, p=1, q=1, X=H,

Y=CO, Z=N and

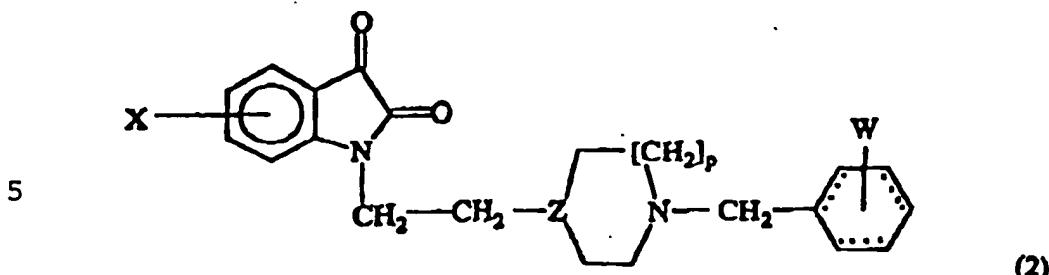


= 4-chlorophenyl are excluded.

35

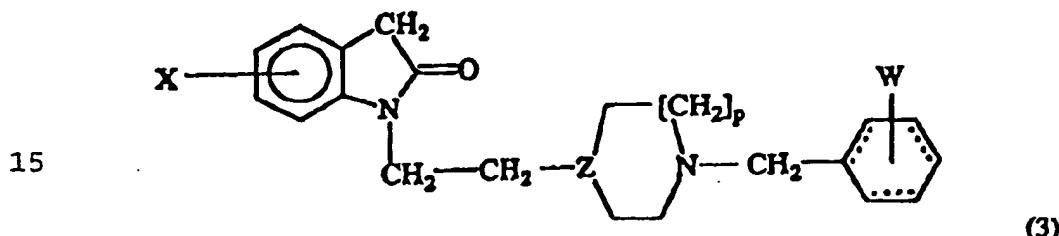
Preferred embodiments of this invention relate to
compounds having the general formula (2)

5



wherein p, X, W and Z are as previously defined above;

10 or to compounds having the general formula (3)



wherein p, X, W and Z are as previously defined above.

20 Throughout the specification and the appended claims, a given chemical formula or name shall encompass all stereo and optical isomers and racemates thereof where such isomers exist, as well as pharmaceutically acceptable acid addition salts thereof and solvates thereof such as for instance hydrates.

25

The following definitions shall apply throughout the specification and the appended claims.

30 Unless otherwise stated or indicated, the term "lower alkyl" denotes a straight or branched alkyl group having from 1 to 6 carbon atoms. Examples of said lower alkyl include methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, sec-butyl, t-butyl and straight- and branched-chain pentyl and hexyl.

35

Unless otherwise stated or indicated, the term

5 "cycloalkyl" denotes a cyclic alkyl group having a ring size from C₃ to C₇, optionally additionally substituted by lower alkyl. Examples of said cycloalkyl include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, methylcyclohexyl and cycloheptyl.

10 Unless otherwise stated or indicated, the term "cycloalkenyl" denotes a cyclic alkenyl group having a ring size from C₃ to C₇, optionally additionally substituted by lower alkyl. Examples of said cycloalkenyl include cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, methylcyclohexenyl and cycloheptenyl.

15 Unless otherwise stated or indicated, the term "aryloxy" denotes a phenoxy group in which the phenyl ring is optionally further substituted by lower alkyl, lower alkoxy or halogen.

20 Unless otherwise stated or indicated, the term "lower alkoxy" denotes a straight or branched alkoxy group having from 1 to 6 carbon atoms. Examples of said lower alkoxy include methoxy, ethoxy, n-propoxy, iso-propoxy, n-butoxy, iso-butoxy, sec-butoxy, t-butoxy and straight- and branched-chain pentoxy and hexoxy.

25 Unless otherwise stated or indicated, the term "halogen" shall mean fluorine, chlorine, bromine or iodine.

30 Unless otherwise stated or indicated, the term "aryl" denotes a phenyl, furyl or thienyl group in which the ring is optionally further substituted by lower alkyl, lower alkoxy or halogen.

35 Unless otherwise stated or indicated, the term "bicycloalkyl" denotes a bicyclic alkyl group having a

siz from C₆ to C₉, optionally additionally substituted by lower alkyl. Examples of said bicycloalkyl include bicyclo[2.2.1]heptyl, bicyclo[2.2.2]octyl and bicyclo[2.2.3]nonyl.

5

Unless otherwise stated or indicated, the term "cyclic acetal" denotes a cyclic acetal group having a ring size from C₅ to C₇. Examples of said cyclic acetal include 1,3-dioxolanyl and 1,3-dioxanyl.

10

Preferred compounds according to the invention are those of general formula (2) or general formula (3) in which:

p is 1,

15

W is hydrogen or F, especially 4-F,
X is lower alkyl, especially methyl or ethyl, lower alkoxy, especially methoxy or ethoxy, cycloalkyl, especially C₅ to C₇ cycloalkyl, F, aryl, especially phenyl, or NR₁R₂, especially 1-pyrrolidinyl or 1-

20

piperidinyl. More preferred compounds according to the invention are those of general formula (2) or general formula (3) in which the X substituent is at the 5-position.

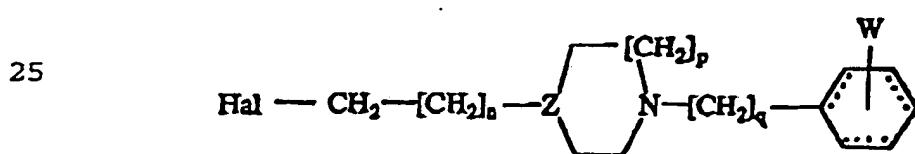
25

Among the most preferred compounds of formula (1) according to the present invention are

- 1,3-Dihydro-5-methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one,
- 5-Cyclohexyl-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one,
- 1,3-Dihydro-1-[2-[4-[(4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-5-methyl-2H-indol-2-one,
- 5-Cyclohexyl-1,3-dihydro-1-[2-[4-[4-fluoro-phenyl)methyl]-1-piperazinyl]ethyl]-2H-indol-2-one,
- 5-Methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione,
- 1-[2-[4-[(4-Fluor ph ny1)methyl]-1-

- piperazinyl]ethyl]-5-methyl-1H-indole-2,3-dione,
 - 5-Cyclohexyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione,
 - 5-Fluoro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione,
 5 - 1,3-Dihydro-5-fluoro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one,
 - 1,3-Dihydro-5-phenyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one,
 10 - 1,3-Dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-5-(1-piperidinyl)-2H-indol-2-one,
 - 5-Cyclohexyl-1,3-dihydro-1-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-2H-indol-2-one
 15 and pharmaceutically acceptable acid addition salts or solvates thereof.

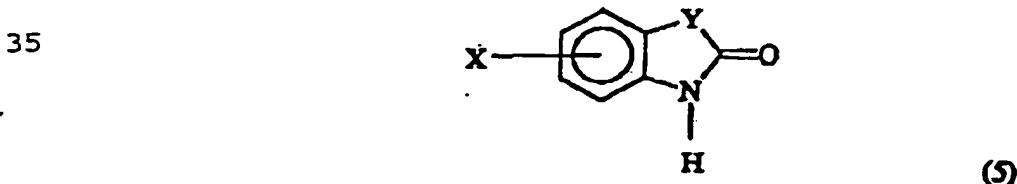
The present invention also relates to processes for preparing the compound having formula (1). Said compound may be prepared by
 20 (a) reacting a compound of the general formula (4) or an acid addition salt thereof



(4)

30 wherein Z, W, n, p and q are as defined above and Hal is halogen,

with a compound of the general formula (5)



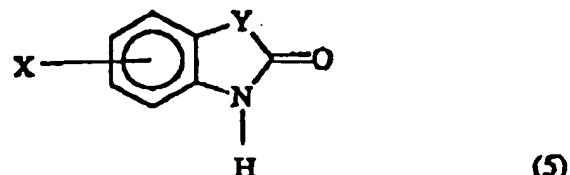
(5)

wherein X and Y are as defined above,

or, in the case where Z=N, by

(b) treating a compound of the general formula (5)

5



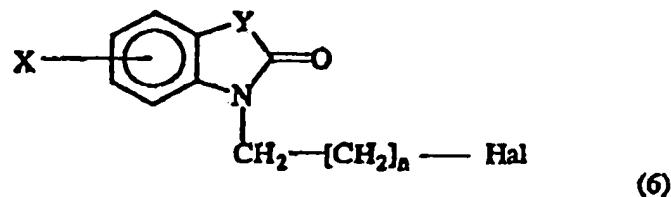
10

wherein X and Y are as defined above,

with a 1,(n+1)-dihaloalkane to obtain a compound of the general formula (6)

15

20

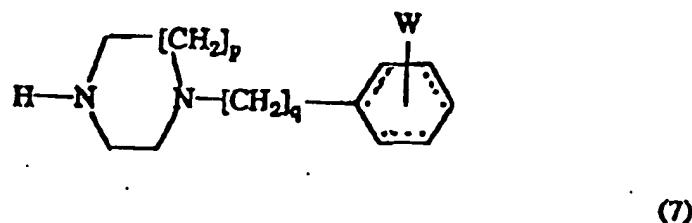


wherein X, Y and n are as defined above and Hal is halogen,

25

and reacting the compound of the general formula (6) with a compound of the general formula (7)

30



wherein W, p and q are as defined above.

35

The process (a) can be achieved, for example, by reacting together a compound of structure (4) or an

acid addition salt thereof with a compound of structure (5) in a suitable solvent such as toluene or 3-methyl-2-butanone or dimethylsulphoxide or dimethylformamide in the presence of a base such as potassium hydroxide or triethylamine or anhydrous potassium carbonate, 5 optionally with the addition of a catalytic amount of potassium iodide. Said reaction should be conducted at a suitable temperature, normally between 0°C and 100°C, optionally in an inert atmosphere. In a preferred variation, 10 a solution of the compound of structure (5) in dimethylformamide at 0°C is treated with a strong base, preferably sodium hydride. After a suitable period of time the compound of structure (4) or an acid addition salt thereof is added to the reaction mixture and the process is then allowed to proceed at ambient 15 temperature or above. The required product (1) may then be isolated and purified and characterised using standard techniques.

The process (b) can be achieved, for example, by 20 treating a compound of structure (5) with a 1,0-dihaloalkane, typically 1-bromo-2-chloroethane, in a suitable solvent such as toluene or 3-methyl-2-butanone or dimethylsulphoxide or dimethylformamide in the presence of a base such as triethylamine or anhydrous 25 potassium carbonate. Such reaction should be conducted at a suitable temperature, normally between 0°C and 100°C, optionally in an inert atmosphere. Some compounds of type (6) are known in the literature. The intermediate (6) may either be isolated and purified 30 and characterised using standard techniques or else may be reacted in a crude form with a compound of structure (7). Such reaction is preferably conducted in a suitable solvent such as dimethylformamide in the presence of a base such as triethylamine or anhydrous 35 potassium carbonate, optionally with the addition of a catalytic amount of potassium iodide. The reaction

should be conducted at a suitable temperature, normally between 0°C and 100°C, optionally in an inert atmosphere. The required product (1) may then be isolated and purified and characterised using standard techniques.

Compounds of structure (4) wherein Hal represents a halogen substituent, preferably either chloro or bromo, are, depending on the nature of the substituent W, either known compounds or compounds which can be prepared using known methods. The application of such methods to the synthesis of compounds of structure (4) will be readily understood by those skilled in the art.

Compounds of structure (5) wherein Y is CO are known as isatins (systematic name 1H-indole-2,3-diones). The isatins of structure (5) are, depending on the nature of the substituent(s) X, either compounds which have been previously described in the literature, or compounds which can be prepared by the straightforward application of known methods. The Sandmeyer procedure (Organic Syntheses, Coll. Vol. I., p 327), in which an aniline, chloral hydrate and hydroxylamine are reacted together to give an intermediate isonitrosoacetanilide which is then cyclised to the isatin on treatment with strong acid, is a particularly useful method.

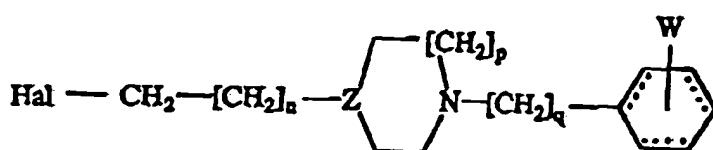
Compounds of structure (5) in which Y is CH₂ are known as oxindoles (systematic name 1,3-dihydro-2H-indol-2-ones). The oxindoles of structure (5) are, depending on the nature of the substituent(s) X, either known compounds or compounds which can be prepared using known methods. The Gassman reaction (P.G. Gassman et al, J.Amer.Chem.Soc., 1974, 96, 5508 and 5512) constitutes a well-known and general synthesis of oxindoles.

12

Compounds of structure (5) wherein Y or Z presents an acetal or cyclic acetal can be prepared from compounds of structure (5) wherein Y is CO by the straightforward application of known methods in a manner that will be readily understood by those skilled in the art.

Thus, the present invention also refers to some new intermediates of formulas (4) and (5), respectively, namely:

10



(4)

wherein Z and Hal are as defined above, n=p=q=1 and

20

W=Me, OMe or F or



=cyclohexyl, with

the proviso that the compound where Z=N and

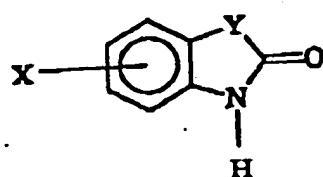
25



=2-methylphenyl is excluded,

and

30



(5)

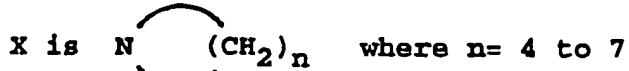
35

wherein

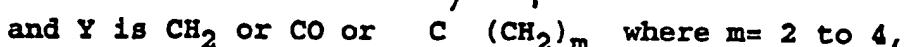
X is cycloalkyl, cycloalkenyl or bicycloalkyl, either

13

optionally further substituted by lower alkyl or



5



10

with the proviso that the compound where X=5-cyclohexyl and Y=CO is excluded.

15 In certain circumstances it is advantageous to prepare oxindoles from the corresponding isatins. This transformation may be achieved using such known methods as:

20 a) catalytic hydrogenation/hydrogenolysis;

b) formation of the corresponding 3-hydrazone followed by reductive elimination under basic conditions (Wolff-Kischner procedure);

25 or

c) formation of the corresponding 3-dithioacetal followed by reduction using Raney nickel or nickel boride.

30 Method (c) represents a preferred process for the conversion of certain isatins (1;Y=CO) or (5;Y=CO) into the corresponding oxindoles (1;Y=CH₂) or (5;Y=CH₂) respectively.

35 The present invention also relates to pharmaceutical formulations containing a compound according to claim 1 as active ingredient and a pharmaceutically acceptable carrier.

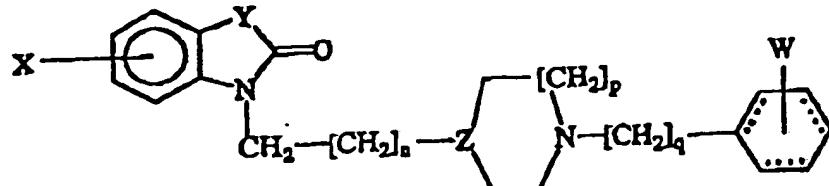
40 Another object of the present invention is a compound

14

according to claim 1 for use in therapy.

Still another object of the present invention is the use of a compound having the general formula (1)

5



10

(I)

wherein:

n is 1, 2 or 3;

15

p is 1 or 2;

q is 1 or 2;

20 X represents one or more substituents independently selected from hydrogen, lower alkyl, aryl, aryloxy, CN, lower alkoxy, halogen, hydroxy, nitro, trifluoromethyl, alkylsulphonamido,

NHCOR where R is lower alkyl or aryl,

25 NR₁R₂ where R₁ and R₂ are independently hydrogen or lower alkyl or together form a ring,

CO₂R where R is lower alkyl,

or cycloalkyl, cycloalkenyl or bicycloalkyl either optionally further substituted by lower alkyl;

30

Y is CO or CR₃R₄ where R₃ and R₄ are independently hydrogen, lower alkyl, lower alkoxy or together form a cyclic acetal;

Z is N or CH;

35

and

represents an optionally substituted

phenyl or cyclohexyl group; wherein

W represents one or more substituents independently selected from hydrogen, lower alkyl, lower alkoxy or halogen;

5

stereo and optical isomers and racemates thereof where such isomers exist, as well as pharmaceutically

10 acceptable acid addition salts thereof and solvates thereof, for the manufacture of a medicament for the treatment of conditions such as glaucoma and myasthenia gravis and, more particularly, for the prevention or treatment of cognitive dysfunctions which may be
15 associated with ageing or with conditions such as Alzheimer's Disease, Senile and related Dementias, Parkinson's Disease, Down's Syndrome and Huntington's Chorea.

20

Moreover, the present invention relates to a method for the treatment of cholinergic dysfunction whereby a pharmacologically effective amount of a compound according to claim 1 is administered to a host in need
25 of said treatment.

Pharmacology

30

The compounds of general formula (1) of the present invention are useful in the treatment of various cognitive dysfunctions, such as those occurring in Alzheimer's disease. This utility is manifested by the ability of these compounds to inhibit the enzyme acetylcholinesterase.

35

Acetylcholinesterase Inhibition Assay

5 The ability of compounds in general to inhibit the acetylcholinesterase activity of rat brain homogenate was determined using the spectrophotometric method of Ellman et al, Biochem.Pharmacol., 1961, 7, 88. Results are expressed as IC₅₀ nanomolar (i.e. the nanomolar concentration of test compound required to inhibit enzyme activity by 50%).

10 Further the compounds of this invention potentiate cholinergic function in the brain such that when administered to rodents these compounds induce marked cholinergic effects such as tremor. These utilities are further demonstrated by the ability of these
15 compounds to restore cholinergically deficient memory in a delayed non-matched to sample task.

Delayed Non-Matched to Sample Assay

20 Rats were trained on a delayed non-matched to sample task similar to that described by Murray et al, Psychopharmacology, 1991, 105, 134-136. Scopolamine, an anticholinergic that is known to cause memory impairment, induces an impairment in performance of this task. This impairment is reversed by compounds of
25 the type described in the present invention.

Pharmaceutical formulations

30 The administration in the novel method of treatment of this invention may conveniently be oral, rectal, or parenteral at a dosage level of, for example, about 0.0001 to 10 mg/kg, preferably about 0.001 to 1.0 mg/kg and especially about 0.01 to 0.2 mg/kg and may be
35 administered on a regimen of 1 to 4 doses or treatments per day. The dose will depend on the route of administration, a preferred route being by oral

5 administration. It will be appreciated that the severity of the disease, the age of the patient and other factors normally considered by the attending physician will influence the individual regimen and dosage most appropriate for a particular patient.

10 The pharmaceutical formulations comprising the compound of this invention may conveniently be tablets, pills, capsules, syrups, powders or granules for oral administration; sterile parenteral solutions or suspensions for parenteral administration; or as suppositories for rectal administration.

15 To produce pharmaceutical formulations containing a compound according to the present invention in the form of dosage units for oral application the active substance may be admixed with an adjuvant/a carrier e.g. lactose, saccharose, sorbitol, mannitol, starches such as potato starch, corn starch or amylopectin, cellulose derivatives, a binder such as gelatine or polyvinylpyrrolidone, and a lubricant such as magnesium stearate, calcium stearate, polyethylene glycol, waxes, paraffin, and the like, and then compressed into tablets. If coated tablets are required, the cores, 20 prepared as described above, may be coated with a concentrated sugar solution which may contain e.g. gum arabic, gelatine, talcum, titanium dioxide, and the like. Alternatively, the tablet can be coated with a polymer known to the man skilled in the art, dissolved 25 in a readily volatile organic solvent or mixture of organic solvents. Dyestuffs may be added to these coatings in order to readily distinguish between tablets containing different active substances or different amounts of the active compounds.

30

35 For the preparation of soft gelatine capsules, the active substance may be admixed with e.g. a veg table

oil or polyethylene glycol. Hard gelatin capsules may contain granules of the active substance using either the above-mentioned excipients for tablets e.g. lactose, saccharose, sorbitol, mannitol, starches (e.g. 5 potato starch, corn starch or amylopectin), cellulose derivatives or gelatine. Also liquids or semisolids of the drug can be filled into hard gelatine capsules.

Dosage units for rectal application can be solutions or 10 suspensions or can be prepared in the form of suppositories comprising the active substance in admixture with a neutral fatty base, or gelatine rectal capsules comprising the active substance in admixture with vegetable oil or paraffin oil.

15 Liquid preparations for oral application may be in the form of syrups or suspensions, for example solutions containing from about 0.02% to about 20% by weight of the active substance herein described, the balance 20 being sugar and mixture of ethanol, water, glycerol and propylene glycol. Optionally such liquid preparations may contain colouring agents, flavouring agents, saccharine and carboxymethylcellulose as a thickening agent or other excipients known to the man in the art.

25 Solutions for parenteral applications by injection can be prepared in an aqueous solution of a water-soluble pharmaceutically acceptable salt of the active substance, preferably in a concentration of from about 30 0.5% to about 10% by weight. These solutions may also contain stabilizing agents and/or buffering agents and may conveniently be provided in various dosage unit ampoules.

EXAMPLE 1

5-(1-Methylethyl)-1H-indole-2,3-dione

4-(1-Methylethyl)-aniline (6.75 g) was dissolved in water (30 ml) containing concentrated hydrochloric acid (4.4 ml). Hydroxylamine hydrochloride (16.9 g) in water (48 ml) and sodium sulphate decahydrate (100 g) in water (120 ml) were added, followed by chloral hydrate (16.5 g) in ethanol (180 ml). The reaction mixture was heated under reflux for 3 hours, then poured into water. The solid isonitroso-acetanilide intermediate was collected by filtration, washed and dried. This material was cooled in an ice-salt bath and concentrated sulphuric acid (48 ml) was added dropwise with stirring. After addition was complete the mixture was warmed to 80°C for 20 minutes and then poured onto crushed ice. The resulting red solid was collected by filtration, washed, dried and then recrystallised from toluene - light petroleum to give the title compound, m.p. 127-129°C.

m/z 207 ($M + NH_4^+$) and 190 ($M + H^+$).
 1H Nmr ($CDCl_3$) 1.16 (6H, d), 2.95 (1H, septuplet), 6.9 (1H, d), 7.45 (1H, dd), 7.5 (1H, d) and 9.0 (1H, br s).

EXAMPLE 2

5-Tetradecyl-1H-indole-2,3-dione

Following the method of Example 1 and starting from 4-tetradecylaniline, the title compound was obtained.

M.p. 87-89°C.

m/z 361 ($M + NH_4^+$) and 344 ($M + H^+$).

30

EXAMPLE 3

5-Cyclohexyl-1,3-dihydro-2H-indol-2-one

5-Cyclohexyl-1H-indole-2,3-dione (3.4 g) in methanol (100 ml) was treated with 1,2-ethanedithiol (1.5 g) and boron trifluoride diethyl etherate (2 ml). The mixture was stirred at room temperature overnight and then evaporated to dryness under reduced pressure. The

residue was purified by flash chromatography to yield the corresponding dithioacetal. This material in ethanol (100 ml) was treated with Raney nickel (50% slurry in water, 40 g) and the mixture was heated under reflux overnight. The mixture was filtered through Celite and the residues washed thoroughly with ethanol. The combined filtrates were evaporated to give the title compound as a white solid (2.9 g, 88%), m.p. 153-155°C.

10 ^1H Nmr (d_6 -DMSO) 1.2-1.5 (5H, m), 1.7-2.0 (5H, m), 2.5 (1H, m), 3.5 (2H, s), 6.8 (1H, d), 7.08 (1H, dd) and 7.15 (1H, d) ppm.

EXAMPLE 4

15 5-Ethyl-1,3-dihydro-2H-indol-2-one
The title compound was prepared from 5-ethyl-1H-indole-2,3-dione following the method of Example 3.
M.p. 136-137°C.
10 ^1H Nmr (CDCl_3) 1.25 (3H, t), 2.6 (2H, q), 3.55 (2H, s), 6.85 (1H, d), 7.05 (1H, dd), 7.1 (1H, d) and 8.9 (1H, br s) ppm.

EXAMPLE 5

25 1-(2-Chloroethyl)-4-[(2-methoxyphenyl)methyl]piperazine
2-Methoxybenzyl chloride (16 g) and 1-(2-hydroxyethyl)piperazine (13 g) in ethanol (50 ml) were heated under reflux for 4 hours. The solvent was removed under vacuum and the resulting oil was passed through a pad of silica gel eluting with 10% methanol-ammonia in dichloromethane to give 1-(2-hydroxyethyl)-4-[(2-methoxyphenyl)methyl]piperazine as a colourless oil (80%), ^{13}C nmr (CDCl_3) 157.4, 130.3, 127.8, 125.2, 119.8, 110.0, 59.5, 57.6, 55.2, 54.8, 52.7 and 52.4 ppm. This material (15 g) was treated at 0°C with thionyl chloride (15 ml). The mixture was then heated at reflux for 2 hours. Toluene was added and the mixture was evaporated under reduced pressure. The

resulting solid was collected and washed thoroughly to give the dihydrochloride of the title compound as a white solid, m.p. 276-279°C (dec.).

5 Found: C, 48.1; H, 6.8; N, 8.0. $C_{14}H_{21}ClN_2O \cdot 2HCl$.
 0.5H₂O requires C, 47.95; H, 6.9; N, 8.0%.

10 This solid was suspended in dichloromethane and extracted twice with 1N sodium hydroxide solution. The organic phase was then washed with water, dried, and evaporated to give 1-(2-chloroethyl)-4-[(2-methoxyphenyl)methyl]piperazine as an oil.

15 ^{13}C Nmr (CDCl₃) 157.6, 130.3, 127.9, 125.5, 120.0, 110.2, 59.6, 55.6, 55.1, 52.9, 52.6 and 40.7 ppm.

20 The following compounds of Examples 6 to 12 were prepared in an analogous manner to that of Example 5 starting from 1-(2-hydroxyethyl)piperazine and the appropriate chloride.

EXAMPLE 6

1-(2-Chloroethyl)-4-[(3-methoxyphenyl)methyl]piperazine

25 ^{13}C Nmr (CDCl₃) 159.4, 139.5, 128.9, 121.2, 114.3, 112.2, 62.6, 59.6, 54.9, 52.9, 52.7 and 40.7 ppm.
Dihydrochloride, m.p. 282-289°C (dec.).

Found: C, 48.1; H, 6.65; N, 7.9. $C_{14}H_{21}ClN_2O \cdot 2HCl$.
 0.5H₂O requires C, 47.95; H, 6.9; N, 8.0%.

30

EXAMPLE 7

1-(2-Chloroethyl)-4-[(3-methylphenyl)methyl]piperazine

35 ^{13}C Nmr (d₆-DMSO) 137.9, 137.0, 129.3, 128.0, 127.4, 125.7, 61.9, 59.1, 52.4, 52.4, 41.3 and 20.8 ppm.
Dihydrochloride - Found: C, 50.6; H, 7.1; N, 8.3.

$C_{14}H_{21}ClN_2 \cdot 2HCl$. 0.5H₂O requires C, 50.2; H, 7.2; N, 8.4%.

EXAMPLE 8**1-(2-Chloroethyl)-4-[(4-fluorophenyl)methyl]piperazine**

5 ^{13}C Nmr (d_6 -DMSO) 162.9 and 159.34 (d, J 241 Hz),
134.20 and 134.15 (d, J 3.4 Hz), 130.46 and 130.34 (d,
J 8.1 Hz), 114.81 and 114.50 (d, J 21 Hz), 61.0, 59.0,
52.4, 52.3 and 41.3 ppm.

Dihydrochloride, m.p. 253-256°C (dec.).

10 Found: C, 47.4; H, 5.9; N, 8.5; F, 5.8. $\text{C}_{13}\text{H}_{18}\text{ClFN}_2$.
2HCl requires C, 47.4; H, 6.1; N, 8.5; F, 5.8%.

EXAMPLE 9**1-(2-Chloroethyl)-4-(cyclohexylmethyl)piperazine**

15 ^{13}C Nmr (d_6 -DMSO) 64.7, 59.1, 53.0, 52.5, 41.4, 34.3,
31.1, 26.3 and 25.4 ppm.

EXAMPLE 10**1-(2-Chloroethyl)-4-(2-phenylethyl)piperazine**

20 ^{13}C Nmr (CDCl_3) 140.1, 128.5, 128.2., 125.9, 60.3,
59.6, 53.0, 52.8, 40.7 and 33.4 ppm.

EXAMPLE 11**1-(2-Chloroethyl)-4-[(3-fluorophenyl)methyl]piperazine**

25 ^1H Nmr (CDCl_3) 2.3-2.6 (8H, m), 2.7 (2H, t), 3.5 (2H,
s), 3.55 (2H, t), 6.9 (1H, m), 7.1 (2H, m) and 7.2-7.3
(1H, m) ppm.

EXAMPLE 12**1-(2-Chloroethyl)-4-[(2-fluorophenyl)methyl]piperazine**

30 ^{13}C Nmr (CDCl_3) 162.3 and 158.7 (d), 130.6 (d), 128.8
(d), 123.9 (d), 123.0 (d), 114.5 and 114.2 (d), 64.0,
54.2, 52.3, 51.8 and 40.2 ppm.

Dihydrochloride - Found: C, 46.1; H, 6.2; N, 8.0.

35 $\text{C}_{13}\text{H}_{18}\text{ClFN}_2$. 2HCl. 0.5H₂O requires C, 46.1; H, 6.25; N,
8.3%.

EXAMPLE 13

5-Methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

5-Methyl-1H-indole-2,3-dione (12.85 g) in dry DMF (50 ml) at 0 to 5°C was treated with sodium hydride (80% dispersion in mineral oil, 2.53 g). The mixture was allowed to warm to room temperature and after a further 10 minutes 1-(2-chloroethyl)-4-(phenylmethyl)piperazine (20 g) in dry DMF (70 ml) was added. The mixture was heated at 70°C for 3 hours and then evaporated under reduced pressure. The residue was subjected to flash chromatography on silica gel to afford the title compound (17.75 g). Treatment with ethanolic HCl then gave 5-methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione dihydrochloride (16.8 g), m.p. 270-275°C (dec.).
¹H Nmr (d₆-DMSO) 2.4 (3H, s), 3.3-3.9 (10H, m), 4.2 (2H, br s), 4.45 (2H, br s), 7.3 (1H, d), 7.45-7.6 (5H, m) and 7.75 (2H, m) ppm.

20

EXAMPLE 14

5-Cyclohexyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

5-Cyclohexyl-1H-indole-2,3-dione (3.45 g) in dry DMF at 0°C was treated with sodium hydride (80% dispersion in mineral oil, 550 mg). The mixture was allowed to warm to room temperature and 1-(2-chloroethyl)-4-(phenylmethyl) piperazine (3.9 g) in dry DMF (25 ml) was added. The reaction mixture was then heated in an oil bath at 70°C for 2 hours. The mixture was evaporated under reduced pressure and the residue passed through a pad of silica gel to yield the title compound as a red oil (4.2 g, 65%).

¹³C Nmr (CDCl₃) 183.8, 158.3, 148.9, 143.7, 137.9, 136.8, 129.0, 128.1, 126.9, 123.4, 117.6, 110.0, 62.8, 54.6, 53.1, 52.8, 43.6, 37.7, 34.2, 26.5 and 25.7 ppm. This oil (4 g) in ethanol (50 ml) was treated with

thanolic HCl to give 5-cyclohexyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione dihydrochloride as an orange solid, m.p. 251-254°C (dec.)

5

The compounds of Examples 15 to 21 were prepared in an analogous manner to Examples 13 and 14, starting from 1-(2-chloroethyl)-4-(phenylmethyl)piperazine and the appropriately substituted 1H-indole-2,3-dione.

10

EXAMPLE 15

5-Butyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

15 ^1H Nmr (CDCl_3) 0.85 (3H, t), 1.25 (2H, m), 1.5 (2H, m), 2.4-2.9 (12H, m), 3.65 (2H, s), 3.75 (2H, t), 6.8 (1H, d) and 7.2-7.4 (7H, m) ppm.

Dihydrochloride, m.p. 217-220°C (dec.).

Found: C, 60.7; H, 7.0; N, 8.7. $\text{C}_{25}\text{H}_{31}\text{N}_3\text{O}_2 \cdot 2\text{HCl} \cdot \text{H}_2\text{O}$ requires C, 60.5; H, 7.1; N, 8.5%.

20

EXAMPLE 16

5-(1-Methylethyl)-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

m/z 391 (M^+), 189 and 91.

25 Dihydrochloride, m.p. 233-234°C (dec.).

Found: C, 58.7; H, 6.7; N, 8.6. $\text{C}_{24}\text{H}_{29}\text{N}_3\text{O}_2 \cdot 2\text{HCl} \cdot 1.5\text{H}_2\text{O}$ requires C, 58.7; N, 7.0; N, 8.55%.

EXAMPLE 17

30 5-Hexyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

m/z 433 (M^+), 189, 91.

Dihydrochloride, m.p. 223-225°C (dec.).

35 ^1H Nmr ($d_6\text{-DMSO}$) 0.9 (3H, t), 1.35 (6H, br s), 1.6 (2H, m), 2.6 (2H, t), 3.4-3.9 (10H, m), 4.15 (2H, br s), 4.45 (2H, br s), 7.25 (1H, d), 7.45 (1H, d), 7.5-7.6 (4H, m) and 7.7 (2H, m) ppm.

EXAMPLE 18

5-Ethyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

5 m/z 377 (M^+), 189, 91.

Dihydrochloride, m.p. 243-245°C (dec.).

Found: C, 60.9; H, 6.1; N, 9.2. $C_{23}H_{27}N_3O_2 \cdot 2HCl$
requires C, 61.3; H, 6.5; N, 9.3%.

10

EXAMPLE 19

1-[2-[4-(Phenylmethyl)-1-piperazinyl]ethyl]-5-tetradecyl-1H-indole-2,3-dione

M.p. 67-68°C.

5 m/z 545 (M^+), 189, 91.

15 Found: C, 75.5; H, 9.65; N, 7.55. $C_{35}H_{51}N_3O_2 \cdot 0.5H_2O$
requires C, 75.8; H, 9.45; N, 7.6%.

EXAMPLE 20

5-(1-Methylpropyl)-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

M.p. 235-236°C (dec.).

20 1H Nmr (d_6 -DMSO) 0.75 (3H, t), 1.15 (3H, d), 1.5 (2H, m), 2.6 (1H, m), 3.3-3.9 (10H, m), 4.1 (2H, br s), 4.4 (2H, br s), 7.25 (1H, d), 7.35-7.6 (5H, m) and 7.7 (2H, m) ppm.

Found: C, 62.5; H, 6.9; N, 8.55; Cl, 14.5. $C_{25}H_{31}N_3O_2 \cdot 2HCl$ requires C, 62.75; H, 6.95; N, 8.8; Cl, 14.8%.

EXAMPLE 21

5-(1,1-Dimethylethyl)-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

M.p. 241-242°C (dec.).

30 1H Nmr (d_6 -DMSO) 1.3 (9H, s), 3.3-3.9 (10H, m), 4.1 (2H, br s), 4.35 (2H, br s), 7.25 (1H, d), 7.45 (3H, m), 7.55 (1H, d), 7.65 (2H, m) and 7.7 (1H, d) ppm.

EXAMPLE 22**1-[2-[4-(Cyclohexylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride**

1H-Indole-2,3-dione (2.4 g) in dry DMF (8 ml) at 0°C
5 was treated with sodium hydride (80% dispersion in
mineral oil, 500 mg). The mixture was allowed to warm
to room temperature and after 30 minutes 1-(2-
chloroethyl)-4-(cyclohexylmethyl)piperazine (4 g) in
10 dry DMF (8 ml) was added. The mixture was heated at
80°C for 1.5 hours and then evaporated under reduced
pressure. The residue was purified by flash
chromatography on silica gel and then treated with
ethanolic HCl to give 1-[2-[4-(cyclohexylmethyl)-1-
15 piperazinyl]ethyl]-1H-indole-2,3-dione dihydrochloride,
m.p. 256-258°C.
Found: C, 57.9; H, 7.6; N, 9.5. $C_{21}H_{29}N_3O_2 \cdot 2HCl$.
0.5H₂O requires C, 57.7; H, 7.4; N, 9.6%.

20 By following the same procedure as in Example 22 but
starting with the appropriate 4-substituted 1-(2-
chloroethyl)piperazine the products of Examples 23 to
27 were obtained.

25

EXAMPLE 23**1-[2-[4-(2-Phenylethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride**

M.p. 252-254°C (dec.).
Found: C, 59.7; H, 6.2; N, 9.2. $C_{22}H_{25}N_3O_2 \cdot 2HCl$.
30 0.5H₂O requires C, 59.3; H, 6.3; N, 9.4%.

EXAMPLE 24**1-[2-[4-[(2-Methoxyphenyl)methyl]-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride**

35 M.p. 224-225°C (dec.).
¹H Nmr (d_6 -DMSO) 3.4-4.0 (10H, m), 3.95 (3H, s), 4.25
(2H, br s), 4.4 (2H, br s), 7.1 (1H, t), 7.2 (2H, m),

7.4 (1H, d), 7.55 (1H, t), 7.65 (1H, d) and 7.75 (2H, m) ppm.

EXAMPLE 25

- 5 1-[2-[4-[(2-Methoxyphenyl)methyl]-1-piperazinyl]ethyl]-
5-(1-methylethyl)-1H-indole-2,3-dione Dihydrochloride
M.p. 215-220°C (dec.).
m/z 422 (M + H⁺).
Found: C, 59.9; H, 6.8; N, 8.4. C₂₅H₃₁N₃O₃. 2HCl.
10 0.5H₂O requires C, 59.6; H, 6.8; N, 8.35%.

EXAMPLE 26

- 15 1-[2-[4-[(3-Methoxyphenyl)methyl]-1-piperazinyl]ethyl]-
1H-indole-2,3-dione Dihydrochloride
M.p. 241-244°C (dec.).
m/z 380 (M + H⁺).
Found: C, 58.0; H, 6.0; N, 9.1. C₂₂H₂₅N₃O₃. 2HCl
requires C, 58.4; H, 6.0; N, 9.3%.

20

EXAMPLE 27

- 1-[2-[4-[(3-Methylphenyl)methyl]-1-piperazinyl]ethyl]-
1H-indole-2,3-dione Dihydrochloride
M.p. 242-245°C (dec.).
m/z 364 (M + H⁺)
25 ¹H Nmr (d₆-DMSO) 2.4 (3H, s), 3.35-4.05 (10H, m), 4.25
(2H, br s), 4.45 (2H, br s), 7.25 (1H, t), 7.3-7.45
(3H, m), 7.55 (1H, d), 7.6 (1H, d), 7.65 (1H, d) and
7.75 (1H, t) ppm.
Found: C, 59.4; H, 6.2; N, 9.4. C₂₂H₂₅N₃O₂. 2HCl. 0.5H₂O
30 requires C, 59.3; H, 6.3; N, 9.4%.

EXAMPLE 28

- 1-[2-[4-[(4-Fluorophenyl)methyl]-1-piperazinyl]ethyl]-
1H-indole-2,3-dione
35 1H-Indole-2,3-dione (2.9 g) in dry DMF (5 ml) at 0°C
was treated with sodium hydride (80% dispersion in
mineral oil, 600 mg). The mixture was warmed to 40°C

and after 45 minutes a solution of 1-(2-chloroethyl)-4-[
5 (4-fluorophenyl)methyl]piperazine (5.1 g) in dry DMF
(8 ml) was added. The reaction mixture was heated at
80°C for 5 hours and then evaporated under reduced
pressure. The residue was recrystallised twice to give
1-[2-[4-[4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-
1H-indole-2,3-dione, m.p. 146-147°C.
10 ^{13}C Nmr (d_6 -DMSO) 183.4, 162.9 and 159.3 (d), 157.9,
150.7, 138.1, 134.2 (d), 130.4 (d), 124.3, 123.0,
117.3, 114.8 and 114.5 (d), 110.9, 61.0, 54.2, 52.6,
52.4 and 37.2 ppm.
Found: C, 68.4; H, 6.3; N, 11.4. $\text{C}_{21}\text{H}_{22}\text{FN}_3\text{O}_2$ requires
C, 68.65; H, 6.0; N, 11.4%.

15 Following the same general method as in Example 28 and
using the appropriately substituted starting materials,
the compounds of Examples 29 to 32 were prepared.

EXAMPLE 29

20 1-[2-[4-[4-Chlorophenyl)methyl]-1-piperazinyl]ethyl]-
1H-indole-2,3-dione
M.p. 126-128°C (dec.).
13C Nmr (d_6 -DMSO) 183.4, 157.9, 150.7, 138.1, 137.1,
131.3, 130.4, 128.0, 124.3, 123.0, 117.3, 110.9, 60.9,
25 54.1, 52.5, 52.4 and 37.2 ppm.

EXAMPLE 30

30 1-[2-[4-[4-Fluorophenyl)methyl]-1-piperazinyl]ethyl]-
5-methyl-1H-indole-2,3-dione
 ^{13}C Nmr (CDCl_3) 183.7, 163.6 and 160.0 (d), 158.3,
148.7, 138.6, 133.6 (d), 133.3, 130.5 (d), 125.6,
117.6, 115.0 and 114.7 (d), 110.1, 62.0, 54.5, 53.1,
52.9, 37.7 and 20.5 ppm.
Dihydrochloride, m.p. 238-240°C (dec.).
35 Found: C, 57.1; H, 5.7; N, 9.2; $\text{C}_{22}\text{H}_{24}\text{FN}_3\text{O}_2 \cdot 2\text{HCl}$.
0.5H₂O requires C, 57.0; H, 5.9; N, 9.1%.

EXAMPLE 31

1-[2-[4-[(2-Fluorophenyl)methyl]-1-piperazinyl]ethyl]-5-methyl-1H-indole-2,3-dione

M.p. 104-106°C.

5 ^1H Nmr (CDCl_3) 2.25 (3H, s), 2.3-2.6 (10H, m), 3.5 (2H, s), 3.7 (2H, t), 6.75 (1H, d) and 6.9-7.4 (6H, m) ppm.
Dihydrochloride, m.p. 240-246°C (dec.).

Found: C, 57.3; H, 5.6; N, 8.9. $\text{C}_{22}\text{H}_{24}\text{FN}_3\text{O}_2 \cdot 2\text{HCl}$. 0.5 H_2O requires C, 57.0; H, 5.9%; N, 9.1%.

10

EXAMPLE 32

1-[2-[4-[(3-Fluorophenyl)methyl]-1-piperazinyl]ethyl]-5-methyl-1H-indole-2,3-dione

15 ^{13}C Nmr (CDCl_3) 183.7, 164.6 and 161.0 (d), 158.3, 148.7, 140.8 (d), 138.6, 133.3, 129.5 (d) 125.6, 124.5 (d), 117.6, 115.8 and 115.5 (d), 114.0 and 113.7 (d), 110.1, 62.2, 54.6, 53.1, 52.9, 37.8 and 20.6 ppm.
Dihydrochloride, m.p. 237-240°C (dec.).

20

EXAMPLE 33

4,7-Dimethyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

4,7-Dimethyl-1H-indole-2,3-dione (700 mg) in dry DMF (10 ml) was cooled to 0°C and sodium hydride (80% dispersion in mineral oil, 120 mg) was added. After 30 minutes at 0°C 1-(2-chloroethyl)-4-(phenylmethyl)piperazine (1 g) in dry DMF (5 ml) was added. The mixture was heated to 80°C for 2 hours and then evaporated under reduced pressure. The residue was subjected to flash chromatography and then treated with ethanolic HCl to give 4,7-dimethyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione dihydrochloride, m.p. 223-227°C (dec.).
 m/z 377 ($M + \text{H}^+$).

35 ^1H Nmr (d_6 -DMSO) 2.5 and 2.55 (each 3H, s), 3.3-4.0 (10H, m), 4.3 (2H, t), 4.45 (2H, br s), 6.9 and 7.4 (ach 1H, d), 7.5 (3H, m) and 7.7 (2H, m) ppm.

Starting from the appropriately substituted 1H-indole-2,3-dione and following the method of Example 33 the compounds of Examples 34 to 42 were prepared.

5

EXAMPLE 34**4-Methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride**

M.p. 228-230°C (dec.).

 m/z 363 (M^+), 189 and 91.10 Found: C, 59.0; H, 6.1; N, 9.5. $C_{22}H_{25}N_3O_2 \cdot 2HCl$.
0.5H₂O requires C, 59.3; H, 6.3; N, 9.4%.**EXAMPLE 35****5-Chloro-7-methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione** ^{13}C Nmr (d_6 -DMSO) 199.3, 169.6, 150.6, 138.3, 135.4, 130.5, 129.8, 129.1, 128.3, 127.1, 120.9, 120.4, 62.3, 57.9, 52.8, 52.7, 44.0 and 20.0 ppm.**Dihydrochloride**, m.p. 241-243°C (dec.).20 m/z 399 and 397 (M^+), 189 and 91.**EXAMPLE 36****5-Chloro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride**

25 M.p. 240-243°C (dec.).

EXAMPLE 37**5-Iodo-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride**

30 M.p. 226-229°C (dec.).

 1H Nmr (d_6 -DMSO) 3.3-4.0 (10H, m), 4.2 (2H, br s), 4.45 (2H, br s), 7.3 (1H, d), 7.5 (3H, m), 7.7 (2H, m), 7.9 (1H, d) and 8.05 (1H, dd) ppm.

35

EXAMPLE 38**4,7-Dichloro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride**

M.p. 248-252°C (dec.).

¹³C Nmr (d₆-DMSO) 177.8, 158.6, 145.7, 139.5, 131.4, 130.3, 129.5, 129.1, 128.7, 125.7, 117.8, 114.2 and 35.4 ppm.

5

EXAMPLE 39

5-Nitro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

¹H Nmr (d₆-DMSO) 2.4-2.6 (8H, m), 2.7 (2H, t), 3.5 (2H, t), 3.55 (2H, s), 7.0 (1H, d), 7.3-7.5 (5H, m), 8.2 (1H, dd) and 8.6 (1H, d) ppm.

Dihydrochloride, m.p. 240-245°C (dec.).

EXAMPLE 40

15 5-Methoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

¹H Nmr (d₆-DMSO) 2.3 (4H, br s), 2.4-2.6 (6H, m), 3.4 (2H, s), 3.7-3.8 (5H, m), 7.15-7.2 (2H, m) and 7.25-7.4 (6H, m) ppm.

20 Dihydrochloride, m.p. 235-245°C (dec.).

Found: C, 58.1; H, 5.9; N, 9.1. C₂₂H₂₅N₃O₃. 2HCl requires C, 58.4; H, 6.0; N, 9.3%.

EXAMPLE 41

25 7-Methoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

M.p. 226-229°C (dec.).

EXAMPLE 42

30 1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-5-trifluoromethyl-1H-indole-2,3-dione

¹H Nmr (CDCl₃) 2.3-2.6 (10H, m), 3.4 (2H, s), 3.8 (2H, t), 7.0 (1H, d), 7.25 (5H, br s) and 7.8 (2H, m) ppm.

Dihydrochloride, m.p. 235-239°C (dec.).

EXAMPLE 43**5-Methyl-1-[3-[4-(phenylmethyl)-1-piperazinyl]propyl]-1H-indole-2,3-dione**

5-Methyl-1H-indole-2,3-dione (1.54 g) in dry DMF (10 ml) at 0°C was treated with sodium hydride (80% dispersion in mineral oil, 300 mg). The mixture was allowed to warm to room temperature and after a further 10 minutes 1-(3-chloropropyl)-4-(phenylmethyl)piperazine (2.53 g) in dry DMF (10 ml) was added. The mixture was heated at 80°C for 3 hours and then evaporated under reduced pressure. The residue was purified by flash chromatography to give the title compound.

15 ^1H Nmr (CDCl_3) 1.85 (2H, m), 2.3 (3H, s), 2.3-2.5 (10H, m), 3.5 (2H, s), 3.75 (2H, t), 6.9 (1H, d), and 7.2-7.4 (7H, m) ppm.

Treatment with ethanolic HCl gave 5-methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]propyl]-1H-indole-2,3-dione dihydrochloride, m.p. 256-261°C (dec.).

20

EXAMPLE 44**1-[3-[4-(Phenylmethyl)-1-piperazinyl]propyl]-1H-indole-2,3-dione Dihydrochloride.**

Following the method of Example 43 but starting with 1H-indole-2,3-dione, there was obtained the title compound.

M.p. 233-236°C (dec.).

30 ^{13}C Nmr (d_6 -DMSO) 183.6, 158.8, 150.6, 138.7, 131.6, 130.0, 129.6, 129.3, 124.9, 123.8, 117.9, 111.0, 59.0, 53.3, 48.2, 47.6, 36.7 and 21.5 ppm.

EXAMPLE 45**5-Methyl-1-[4-[4-(phenylmethyl)-1-piperazinyl]butyl]-1H-indole-2,3-dione**

35 5-Methyl-1H-indole-2,3-dione (1.6 g) in dry DMF (20 ml) at 0°C was treated with sodium hydride (80% dispersion in mineral oil, 300 mg). After 30 minutes at 0°C, 4-

br mo-1-chlorobutan (6.8 g) was added and the mixture was then heated at 90°C for 2 hours. The mixture was evaporated to dryness under reduced pressure and the residue was treated with 1-benzylpiperazine (1.76 g) in dry DMF (20 ml). The resulting mixture was heated to 90°C for 4 hours and then left to stand at room temperature overnight. The mixture was evaporated to dryness under reduced pressure and the residue was purified by flash chromatography to yield the title compound.

13C Nmr (CDCl₃) 183.9, 158.2, 148.7, 138.7, 137.9, 133.5, 129.2, 128.2, 127.1, 125.8, 117.6, 110.1, 63.0, 57.6, 53.1, 53.0, 39.9, 25.0, 24.0 and 20.7 ppm. Treatment with ethanolic HCl gave 5-methyl-1-[4-[4-(phenylmethyl)-1-piperazinyl]butyl]-1H-indole-2,3-dione dihydrochloride, m.p. 235-238°C (dec.).

EXAMPLE 46

20 1-[3-[4-(Phenylmethyl)-1-(hexahydro-1H-1,4-diazepinyl)]propyl]-1H-indole-2,3-dione

Sodium hydride (80% dispersion in mineral oil, 140 mg) was added to a solution of 1H-indole-2,3-dione (660 mg) in dry DMF (6 ml) at 0°C. The mixture was allowed to warm to room temperature and after 30 minutes a solution of 1-(3-chloropropyl)-4-(phenylmethyl)-hexahydro-1H-1,4-diazepine (1.3 g) in dry DMF (8 ml) was added. The mixture was stirred at room temperature for 1 hour and then at 80°C for 1 hour. The mixture was evaporated to dryness under reduced pressure and the residue was purified by flash chromatography to give the title compound as a red oil (820 mg, 48%).

1H Nmr (d₆-DMSO) 1.6-1.7 (4H, m), 2.4-2.7 (10H, m), 3.5 (2H, s), 3.7 (2H, t) and 7.1-7.7 (9H, m) ppm.

5,6-Dimethoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

- Anhydrous potassium carbonate (2.44 g) was added to a solution of 5,6-dimethoxy-1H-indole-2,3-dione (1.2 g) in dry DMF (5 ml). 2-Bromo-1-chloroethane (4.1 g) was added and the mixture was heated at 70°C for 2 hours.
- 5 The mixture was evaporated to dryness under reduced pressure and the residue was purified by flash chromatography on silica gel. The 1-(2-chloroethyl)-5,6-dimethoxy-1H-indole-2,3-dione thus obtained was dissolved in dry DMF (5 ml) and anhydrous potassium carbonate (2.44 g), potassium iodide (100 mg) and 1-(phenylmethyl)piperazine (3.06 g) were added. The mixture was stirred and heated at 70°C for 2 hours and then evaporated to dryness under reduced pressure. The residue was purified by chromatography to yield a red oil which on treatment with ethanolic HCl afforded 5,6-dimethoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione dihydrochloride (33%), m.p. 205-207°C (dec.).
- 10 Following the general procedure of Example 47 but using the appropriately substituted 1H-indole-2,3-dione, the products of Examples 48 and 49 were prepared.

EXAMPLE 48

- 25 6-Methoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione
M.p. 136-138°C.

EXAMPLE 49

- 30 7-Methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione
 ^{13}C Nmr (d_6 -DMSO) 200.6, 170.6, 151.9, 138.3, 136.5, 132.2, 129.2, 128.4, 127.2, 127.1, 120.1, 117.5, 62.3, 58.1, 52.7, 52.6, 44.2 and 20.3 ppm.
- 35 Dihydrochloride, m.p. 248-249°C (dec.).

EXAMPLE 50**1,3-Dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one**

Sodium hydride (80% dispersion in mineral oil, 250 mg) was added to a solution of 1,3-dihydro-2H-indol-2-one (1.12 g) in dry DMF (5 ml) at 0°C. The mixture was allowed to warm to room temperature and after 50 minutes a solution of 1-(2-chloroethyl)-4-(phenylmethyl)piperazine (2.02 g) in dry DMF (6 ml) was added. The reaction mixture was then heated at 80°C for 2 hours and then evaporated to dryness under reduced pressure. The residue was purified by flash chromatography on silica gel to afford the title compound (1.1 g, 40%) as an oil.

¹³C Nmr (d₆-DMSO) 174.1, 144.2, 138.1, 128.7, 128.0, 127.4, 126.7, 124.6, 124.1, 121.5, 108.3, 62.0, 54.6, 52.7, 52.5, 36.9 and 35.0.

Treatment with ethanolic HCl gave 1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one dihydrochloride, m.p. 253-256°C (dec.).

EXAMPLE 51**1,3-Dihydro-3,3-dimethyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one Dihydrochloride**

Following the general method of Example 50 but starting with 1,3-dihydro-3,3-dimethyl-2H-indol-2-one, there was obtained 1,3-dihydro-3,3-dimethyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one dihydrochloride, m.p. 218-220°C (dec.).

Found: C, 62.0; H, 7.4; N, 9.3. C₂₃H₂₉N₃O. 2HCl. 1.5 H₂O requires C, 62.0; H, 7.2; N, 9.4%.

Starting with the appropriately substituted 1,3-dihydro-2H-indol-2-one and following the general method of Example 50 the compounds of Examples 52 to 54 were prepared.

EXAMPLE 52

1,3-Dihydro-7-methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one Dihydrochloride

M.p. 234-236°C (dec.).

5 m/z 349 (M^+), 189 and 91.

EXAMPLE 53

1,3-Dihydro-5-methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

10 ^1H Nmr (CDCl_3) 2.3 (3H, s), 2.4-2.7 (10H, m), 3.4 (2H, s), 3.55 (2H, s), 3.75 (2H, t), 6.5 (1H, d), 7.0 (2H, m) and 7.2-7.35 (5H, m) ppm.

Dioxalate, m.p. 219-223°C (dec.).

15 Found: C, 57.3; H, 5.9; N, 7.5. $\text{C}_{22}\text{H}_{27}\text{N}_3\text{O}_2 \cdot 2(\text{CO}_2\text{H})_2$. H_2O requires C, 57.0; H, 6.1; N, 7.7%.

EXAMPLE 54

5-Cyclohexyl-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

20 ^1H Nmr (CD_2Cl_2) 1.1-1.4 (5H, m), 1.6-1.8 (5H, m), 2.2-2.55 (11H, m), 3.3 (2H, s), 3.35 (2H, s), 3.65 (2H, t), 6.65 (1H, d), 6.95 (1H, dd), 7.0 (1H, d) and 7.1-7.25 (5H, m) ppm.

Dihydrochloride, m.p. 218-220.5°C (dec.).

25 Found: C, 64.1; H, 7.7; N, 8.0. $\text{C}_{27}\text{H}_{35}\text{N}_3\text{O} \cdot 2\text{HCl} \cdot \text{H}_2\text{O}$ requires C, 63.8; H, 7.7; N, 8.3%.

EXAMPLE 55

1,3-Dihydro-1-[3-[4-(phenylmethyl)-1-

30 piperazinyl]propyl]-2H-indol-2-one Dioxalate

Following the method of Example 43 but starting with 1,3-dihydro-2H-indol-2-one there was obtained the title compound. M.p. 216-217°C.

35 Found: C, 56.9; H, 5.7; N, 7.5. $\text{C}_{22}\text{H}_{27}\text{N}_3\text{O} \cdot 2\text{C}_2\text{H}_2\text{O}_4$. H_2O requires C, 57.0; H, 6.1; N, 7.7%.

Following the method of Example 50 but starting with

the appropriately substituted 1,3-dihydro-2H-indol-2-one, the compounds of Examples 56 to 59 were prepared.

EXAMPLE 56

5-Cyclopentyl-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

5 ^{13}C Nmr(CDCl₃) 174.6, 142.0, 140.1, 137.7, 128.8,
127.8, 126.7, 125.8, 124.3, 123.0, 107.6, 62.7, 54.6,
52.9, 52.6, 45.3, 37.3, 35.5, 34.4 and 25.2 ppm.

10

EXAMPLE 57

1,3-Dihydro-5-(1-methylpropyl)-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

15 ^{13}C Nmr(CDCl₃) 174.5, 142.0, 141.3, 137.5, 128.8,
127.8, 126.7, 125.9, 124.3, 122.8, 107.6, 62.5, 54.5,
52.8, 52.5, 41.0, 37.2, 35.4, 31.0, 21.9 and 12.0 ppm.

EXAMPLE 58

1,3-Dihydro-5-ethyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

20 ^{13}C Nmr(CDCl₃) 174.9, 142.1, 138.3, 137.5, 129.2,
128.2, 127.1, 126.8, 124.6, 124.1, 108.0, 62.8, 54.7,
53.0, 52.8, 37.5, 35.7, 28.4 and 16.0 ppm.

EXAMPLE 59

25 1,3-Dihydro-5-nitro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

M.p. 134 - 136°C.

^{13}C Nmr(CDCl₃) 174.7, 150.3, 143.0, 137.6, 129.2,
128.2, 127.1, 125.1, 125.0, 120.2, 107.9, 62.8, 54.9,

30 53.2, 52.9, 38.1 and 35.2 ppm.

Dioxalate, m.p. 205 - 208°C (dec.)

Following the general method of Example 14 but starting with the appropriately substituted 1H-indole-2,3-dione,

the compounds of Examples 60 to 67 were prepared.

EXAMPLE 60

5-Cyclopentyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

5 ^{13}C Nmr(CDCl₃) 183.9, 158.5, 148.9, 142.4, 137.9,
137.1, 129.2, 128.2, 127.1, 123.9, 117.7, 110.0, 63.0,
54.7, 53.2, 52.9, 45.1, 37.9, 34.5 and 25.3 ppm.
10 Dihydrochloride, m.p. 232 - 240°C (dec.).
 m/z 418 (M + H⁺).

EXAMPLE 61

7-Cyclopentyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

15 ^{13}C Nmr(CDCl₃) 184.0, 160.6, 147.6, 138.0, 137.5,
131.6, 129.1, 128.1, 127.0, 124.1, 123.1, 119.8, 62.9,
55.2, 53.1, 53.0, 41.3, 39.0, 34.9 and 25.5 ppm.

EXAMPLE 62

5-Cycloheptyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

20 ^{13}C Nmr(CDCl₃) 183.8, 158.4, 148.7, 145.7, 137.9,
136.6, 129.1, 128.1, 127.0, 123.4, 117.6, 110.0, 62.9,
54.6, 53.2, 52.9, 46.1, 37.8, 36.6, 27.7 and 26.9 ppm.

EXAMPLE 63

7-Cycloheptyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

30 ^{13}C Nmr(CDCl₃) 184.0, 160.1, 145.8, 137.9, 137.8,
134.6, 129.2, 128.2, 127.0, 124.2, 123.1, 119.5, 63.0,
55.3, 53.2, 53.0, 41.1, 39.2, 36.6, 27.4 and 26.8 ppm.

EXAMPLE 64

35 5-Phenoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride
 M.p. 233 - 236°C (d.c.)

^{m/z} 442 (M + H⁺)

Found: C, 61.8; H, 5.6; N, 8.0. C₂₇H₂₇N₃O₃. 2HCl. 0.5 H₂O requires C, 61.95; H, 5.8; N, 8.0%

5

EXAMPLE 65

5-Cyano-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

M.p. 244 - 246°C (dec.).

^{m/z} 375 (M + H⁺).

10

EXAMPLE 66

5-Fluoro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione Dihydrochloride

^{m/z} 368 (M + H⁺)

Found: C, 57.0; H, 5.6; N, 9.5. C₂₁H₂₂N₃O₂F. 2HCl requires C, 57.3; H, 5.5; N, 9.5%

20

EXAMPLE 67

5-Ethoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

¹³C Nmr(CDCl₃) 183.8, 158.2, 155.5, 144.6, 137.9, 129.0,

25 128.0, 126.8, 125.0, 117.8, 111.2, 109.9, 64.1, 62.8, 54.6, 52.9, 52.7, 37.7 and 14.5 ppm.

EXAMPLE 68

5-Amino-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

1,3-Dihydro-5-nitro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one (200 mg) in ethanol (100 ml) containing 5% palladium on carbon (60 mg) was stirred under an atmosphere of hydrogen at STP for 1 hour. The catalyst was filtered off, the filtrate evaporated to dryness, and the residue purified by flash chromatography on silica gel.

35

40

^{13}C Nmr (CDCl_3) 174.3, 141.8, 137.9, 136.4, 129.1, 128.1, 126.9, 125.8, 113.5, 112.7, 108.6, 62.9, 54.8, 53.1, 52.8, 37.5 and 35.9 ppm.

Trihydrochloride, m.p. 205 - 220°C (dec.).

5

EXAMPLE 69

5-Acetylamino-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

The compound of Example 68 and triethylamine in dry dichloromethane were treated with acetyl chloride.

10 After 2 hours at RT the reaction was worked up and the product purified by flash chromatography on silica gel to afford the title compound, m.p. 145 - 147°C.

15 ^{13}C Nmr (CDCl_3) 174.7, 168.5, 141.0, 137.9, 132.8, 129.2, 128.2, 127.0, 125.2, 119.9, 118.0, 108.2, 62.9, 54.8, 53.2, 52.9, 37.7, 35.9 and 24.3 ppm.

20 Following the general method of Example 50 but starting with the appropriately substituted 1,3-dihydro-2H-indol-2-one and using 1-(2-chloroethyl)-4-[(4-fluorophenyl)methyl]piperazine, the compounds of Examples 70 to 74 were obtained:

EXAMPLE 70

25 1,3-Dihydro-1-[2-[4-[(4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-2H-indol-2-one Dioxalate

M.p. 202 - 205°C (dec.).

m/z 354 (M + H⁺).

Found: C, 55.5; H, 5.3; N, 7.8. $\text{C}_{21}\text{H}_{24}\text{N}_3\text{OF}$. 2 oxalate.

30 0.5H₂O requires C, 55.35; H, 5.4; N, 7.75%

EXAMPLE 71

1,3-Dihydro-1-[2-[4-[(4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-5-methyl-2H-indol-2-one Dioxalate

35 M.p. 206 - 208°C (dec.).

m/z 368 (M + H⁺).

Found: C, 55.6; H, 5.5; N, 7.1. $\text{C}_{22}\text{H}_{26}\text{N}_3\text{OF}$. 2 oxalate.

41

H_2O requires C, 55.2; H, 5.7; N, 7.4%

EXAMPLE 72

- 5 5-Cyclohexyl-1,3-dihydro-1-[2-[4-[(4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-2H-indol-2-one
- 13C Nmr (CDCl₃) 174.6, 163.4 and 159.8 (doublet),
142.0,
133.4, 130.3, 130.2, 125.6, 124.3, 122.8, 114.8 and
10 114.5 (doublet), 107.7, 61.8, 54.6, 52.9, 52.5, 43.9,
37.2, 35.5, 34.4, 26.5 and 26.0 ppm.

EXAMPLE 73

- 15 1,3-Dihydro-5-fluoro-1-[2-[4-[(4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-2H-indol-2-one Dihydrochloride
- M.p. 227-235°C (dec.).
- Found: C, 54.8; H, 5.7; N, 8.8. C₂₁H₂₃F₂N₃O.
2HCl. H_2O requires C, 54.6; H, 5.9; N, 9.1%

20 EXAMPLE 74

- 1,3-Dihydro-5-ethyl-1-[2-[4-[(4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-2H-indol-2-one Dihydrochloride
- M.p. 242-243°C (dec.).
- Found: C, 58.4; H, 6.4; N, 8.6. C₂₃H₂₈N₃OF.
2HCl. H_2O requires C, 58.5; H, 6.8; N, 8.9%

Following the general method of Example 50 but starting with the appropriately substituted 1,3-dihydro-2H-indol-2-one, the compounds of Examples 75 to 88 were obtained.

EXAMPLE 75

- 1,3-Dihydro-5-fluoro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one
- 35 13C Nmr (CDCl₃) 174.5, 160.8 and 157.0 (doublet),
140.5, 137.8, 129.2, 128.2, 127.0, 126.1 and 126.0
(doublet), 114.1 and 113.7 (doublet), 112.7 and 112.3

42

(doubl t), 108.7 and 108.6 (d ublet), 62.9, 54.8, 53.2,
52.9, 37.8 and 35.9 ppm.

Dihydrochloride, m.p. 214-219°C (dec.).

5

EXAMPLE 76

1,3-Dihydro-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-5-trifluoromethyl-2H-indol-2-one
Dihydrochloride

M.p. 233-237°C (dec.).

10

EXAMPLE 77

1,3-Dihydro-7-fluoro-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-2H-indol-2-one Dihydrochloride
M.p. 240-247°C.

15 Found: C, 56.6; H, 6.3; N, 9.5. C₂₁H₂₄FN₃O.
2HCl. H₂O requires C, 56.8; H, 6.3; N, 9.5%

20

EXAMPLE 78

5-Bromo-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-2H-indol-2-one Dihydrochloride
M.p. 260-264°C (dec.).

25

EXAMPLE 79

5-Cyano-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-2H-indol-2-one
¹³C Nmr (CDCl₃) 174.1, 148.3, 137.7, 132.9, 128.9,
127.9, 127.4, 126.8, 125.2, 119.0, 108.6, 104.8, 62.7,
54.7, 53.0, 52.7, 37.7 and 34.8 ppm.
Dihydrochloride, m.p. 247-252°C (dec.).

30

EXAMPLE 80

7-Cycloheptyl-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-2H-indol-2-one
¹³C Nmr (CDCl₃) 175.9, 139.6, 137.9, 132.4, 129.2,
128.1, 127.0, 126.9, 125.3, 122.4, 121.7, 63.0, 55.9,
53.3, 53.0, 40.4, 38.8, 37.2, 35.4, 27.5 and 27.2 ppm.
Dihydrochloride m.p. 210-215°C (d.c.).

43

EXAMPLE 81

5-Cycloheptyl-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one Dihydrochloride
M.p. 212-216°C (dec.).

5

EXAMPLE 82

5-Diethylamino-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one
¹³C Nmr (CDCl₃) 174.5, 144.5, 137.9, 134.7, 129.2,
10 128.2, 127.0, 126.0, 111.9, 111.1, 108.8, 63.0, 54.9,
53.3, 53.0, 44.9, 37.6, 36.4 and 12.5 ppm.
Trihydrochloride, m.p. 188-193°C (dec.).
m/z 406 (M⁺), 189, 91.

15

EXAMPLE 83

1,3-Dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-5-(1-pyrrolidinyl)-2H-indol-2-one
¹³C Nmr (CDCl₃) 174.3, 144.7, 137.9, 134.0, 129.2,
128.2, 127.0, 125.9, 109.8, 109.2, 108.8, 63.0, 54.9,
20 53.4, 52.9, 48.0, 37.6, 36.3 and 25.3 ppm.
Trihydrochloride, m.p. 233-239°C.
m/z 404 (M⁺), 189, 91.
Found: C, 57.7; H, 7.3; N, 10.4. C₂₅H₃₂N₄O.
3HCl. 0.5 H₂O requires C, 57.4; H, 6.9; N, 10.7%

25

EXAMPLE 84

1,3-Dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-5-(1-piperidinyl)-2H-indol-2-one
¹³C Nmr (CDCl₃) 174.6, 148.7, 137.9, 137.5, 129.2,
30 128.1, 127.0, 125.4, 116.2, 115.4, 108.3, 62.9, 54.9,
53.2, 53.0, 52.2, 37.6, 36.1, 26.0 and 24.0 ppm.

EXAMPLE 85

1,3-Dihydro-5-ethoxycarbonyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one
¹³C Nmr (CDCl₃) 175.0, 166.2, 148.5, 137.8, 130.3,
35 129.0, 128.0, 126.9, 125.5, 124.3, 124.2, 107.7, 62.8,

60.6, 54.8, 53.1, 52.8, 37.8, 35.2 and 14.2 ppm.

EXAMPLE 86

- 5 1,3-Dihydro-5-methoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one
 ^{13}C Nmr (CDCl₃) 174.2, 155.3, 137.75, 137.7, 128.9, 127.9, 126.7, 125.6, 111.7, 111.6, 108.2, 62.7, 55.4, 54.6, 53.0, 52.7, 37.4 and 35.8 ppm.

10

EXAMPLE 87

- 1,3-Dihydro-6-methoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one
15 ^{13}C Nmr (CDCl₃) 175.3, 159.7, 145.2, 137.6, 128.8, 128.0, 126.7, 124.5, 116.0, 105.5, 96.1, 62.6, 55.1, 54.6, 52.9, 52.5, 37.1 and 34.7 ppm.

EXAMPLE 88

- 20 1,3-Dihydro-4,5-dimethoxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one
 ^{13}C Nmr (CDCl₃) 173.9, 147.6, 145.5, 138.3, 137.6, 128.7, 127.9, 126.7, 115.6, 111.2, 102.1, 62.6, 59.4, 55.9, 54.5, 53.0, 52.6, 37.4 and 33.8 ppm.

25

5-Benzoylamino-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one

- The compound of Example 68 and triethylamine in dry dichloromethane were treated with benzoyl chloride. After 1 hour at RT the reaction was worked up and the product purified by flash chromatography on silica gel to afford the title compound.

30 ^{13}C Nmr (CDCl₃) 174.8, 165.8, 141.1, 137.8, 134.7, 133.0, 131.8, 129.0, 128.5, 128.0, 127.1, 127.0, 125.1, 120.3, 118.2, 108.2, 62.9, 54.7, 53.1, 52.7, 37.7 and 35.8 ppm.

Dihydrochloride, m.p. 253-256°C (dec.).

EXAMPLE 90**1,3-Dihydro-5-methylsulphonamido-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one**

5 The compound of Example 68 in diethylether was treated with methanesulphonyl chloride. After 2 hours at RT the reaction was worked up and the product purified by flash chromatography on silica gel to yield the title compound.

M.p. 196-198°C.

10 ^{m/z} 428 (M⁺), 189 and 91.

EXAMPLE 91**1,3-Dihydro-5-hydroxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one**

15 The compound of Example 86 in dry dichloromethane at -70°C was treated under an atmosphere of dry nitrogen with boron tribromide (3.5 equivalents). The reaction mixture was allowed to warm to RT, stirred for 2 hours, and then evaporated under reduced pressure.

20 The residue was stirred at RT with methanol for 1hr and then worked up in the usual manner to give the title compound.

¹³C Nmr (CDCl₃) 175.1, 152.5, 137.3, 136.2, 129.3, 128.2, 127.1, 125.8, 113.9, 112.9, 108.5, 62.8, 54.7, 52.9, 52.2, 36.9 and 36.1 ppm.

EXAMPLE 92**1,3-Dihydro-4,5-dihydroxy-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one**

30 The compound of Example 88 was treated by the general method of Example 91 to afford the title compound.

¹³C Nmr (d₆-DMSO) 173.8, 142.1, 141.3, 137.7, 136.8, 128.8, 128.1, 126.9, 113.7, 110.5, 99.0, 61.8, 54.6, 52.5, 52.3, 36.9 and 33.4 ppm.

EXAMPLE 93**5'-Cyclohexyl-spiro[1,3-dioxolane-2,3'-[3H]-indol]-2'(1'H)-one**

5-Cyclohexyl-1H-indole-2,3-dione (1 equivalent),
ethane-1,2-diol (5 equivalents) and p-toluenesulphonic
acid (0.02 equivalents) in dry toluene were heated
under reflux overnight with azeotropic removal of
water. The reaction mixture was cooled, washed with
saturated sodium bicarbonate solution, and then worked
up in the usual manner to afford the title compound.

10 M.p. 178-180°C.

^{13}C Nmr (CDCl_3) 175.8, 143.4, 139.6, 129.9, 124.1,
123.4, 110.5, 102.6, 65.7, 44.1, 34.5, 26.8 and 26.0
ppm.

15

EXAMPLE 94**5'-Phenyl-spiro[1,3-dioxolane-2,3'-[3H]-indol]-2'(1'H)-one**

5'-Bromo-spiro[1,3-dioxolane-2,3'-[3H]indol]-2'(1'H)-one (5.3 g) in dimethoxyethane (130 ml) and ethanol
(33 ml) was treated with phenylboronic acid (7.2 g),
tetakis(triphenylphosphine)palladium (0) (0.5 g),
triethylamine (4.1 ml) and 2M aqueous sodium carbonate
(19.6 ml). The mixture was refluxed overnight, cooled,
and filtered through a pad of silica gel. The filtrate
was evaporated to dryness and the residue crystallised
from ethyl acetate.

M.p. 189-191°C.

m/z 267.

30 ^{13}C Nmr (d_6 -DMSO) 174.4, 142.1, 139.5, 134.6, 129.8,
128.8, 127.0, 126.1, 125.5, 123.0, 110.8, 101.6 and
65.5 ppm.

EXAMPLE 95**35 5'-(Bicyclo[2.2.1]hept-2-yl)-spiro[1,3-dioxolane-2,3'-[3H]-indol]-2'(1'H)-one**

5'-Iodo-spiro[1,3-dioxolane-2,3'-[3H]indol]-2'(1'H)-one

(3.5 g), bicyclo[2.2.1]heptene (1.15 g), piperidine (3.2 g) and bis(triphenylphosphine)palladium (II) acetate (0.35 g) in DMF (5 ml) and formic acid (1.1 ml) were heated and stirred under nitrogen at 60°C for 1 hour. The mixture was cooled, water (50 ml) and ethyl acetate (50 ml) were added, and after 5 minutes the organic layer was separated, washed, dried and evaporated to dryness. The residue was purified by flash chromatography to yield the title compound (60%).
10 M.p. 159-161°C.

EXAMPLE 96

5'-Phenyl-1'-(2-[4-(phenylmethyl)-1-piperazinyl]ethyl)-spiro[1,3-dioxolane-2,3'-(3H)indol]-2'(1'H)-one

15 Following the general method of Example 14, 5'-phenyl-spiro[1,3-dioxolane-2,3'-(3H)indol]-2'(1'H)-one and 1-(2-chloroethyl)-4-(phenylmethyl)piperazine were reacted together to give the title compound.
20 ^{13}C Nmr (d_6 -acetone) 173.9, 144.5, 141.0, 139.6, 136.6, 130.7, 129.7, 129.6, 128.9, 127.9, 127.6, 127.3, 126.2, 124.1, 110.6, 102.7, 66.5, 63.3, 55.6, 54.0, 53.8 and 38.0 ppm.
Dihydrochloride, m.p. 252-254°C (dec.).

25

EXAMPLE 97

5-Phenyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione

The compound of Example 96 in a mixture of tetrahydrofuran (40 ml) and 3M hydrochloric acid (20 ml) was heated under reflux overnight. The mixture was cooled, basified by the addition of saturated aqueous sodium bicarbonate, and extracted with dichloromethane to yield the title compound.
30 ^{13}C Nmr (d_6 -acetone) 184.0, 158.5, 150.9, 139.5, 139.1, 136.7, 136.5, 129.3, 129.2, 128.4, 127.9, 127.1, 126.8, 122.8, 118.6, 111.8, 62.8, 55.0, 54.4, 53.3 and 38.0 ppm.
35

Dihydrochloride, m.p. 262-265°C (d.c.).

EXAMPLE 98

5 5-(Bicyclo[2.2.1]hept-2-yl)-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-1H-indole-2,3-dione

Following the methods of Examples 96 and 97, 5'-
(bicyclo[2.2.1]hept-2-yl)-spiro[1,3-dioxolane-2,3'-
[3H]indol]-2'(1'H)-one was converted into 5'-
(bicyclo[2.2.1]hept-2-yl)-1'-[2-[4-(phenylmethyl)-1-
10 piperazinyl]ethyl]-spiro[1,3-dioxolane-2,3'-[3H]-
indol]-2'(1'H)-one and thence into the title compound.
 ^{13}C Nmr (CDCl_3) 183.9, 158.4, 148.6, 143.4, 137.3,
137.0, 129.3, 128.2, 127.3, 123.5, 117.5, 110.0, 62.7,
54.5, 52.8, 52.7, 46.4, 42.7, 39.0, 37.7, 36.8, 35.9,
15 30.3 and 28.6 ppm.

Dihydrochloride, m.p. 242-245°C (dec.).

EXAMPLE 99

20 1,3-Dihydro-5-phenyl-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-2H-indol-2-one

The compound of Example 97 (400 mg), ethane-1,2-dithiol
(100 mg) and p-toluenesulphonic acid (500 mg) in
glacial acetic acid (10 ml) were stirred at RT
overnight. The mixture was evaporated to dryness. The
25 residue was treated with aqueous sodium bicarbonate and
extracted with dichloromethane. The extracts were
washed, dried and evaporated to give 1,3-dihydro-3,3-
ethylenedithio-5-phenyl-1-[2-[4-(phenylmethyl)-1-
piperazinyl]ethyl]-2H-indol-2-one.

30 To this product (500 mg) in methanol (13 ml) and
tetrahydrofuran (4 ml) was added nickel (II) chloride
hexahydrate (1.6 g). The mixture was cooled to 0°C and
after 5 minutes, sodium borohydride (760 mg) was added.
After a further 30 minutes at 0°C, the mixture was
35 filtered through a pad of C lit. The filtrate was
evaporated to dryness. The residue was dissolved in
methanol (30 ml), 3M hydrochloric acid (20 ml) was

added, and the mixture was heated under reflux for 2 hours. The methanol was removed under reduced pressure and the remaining aqueous solution was basified by the addition of saturated aqueous sodium bicarbonate. The 5 mixture was extracted with dichloromethane. The material thus obtained was purified by flash chromatography to give the title compound.

13C Nmr (d₆-acetone) 175.5, 145.7, 142.3, 140.1, 136.1, 130.2, 130.1, 129.4, 128.1, 128.0, 127.8, 127.5, 126.9, 10 124.4, 109.9, 63.9, 56.3, 54.6, 54.4, 38.7 and 36.4 ppm.

Dihydrochloride, m.p. 256-258°C (dec.).

EXAMPLE 100

15 5-(Bicyclo[2.2.1]hept-2-yl)-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one
Following the general method of Example 99, 5-(bicyclo[2.2.1]hept-2-yl)-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione was converted 20 into 5-(bicyclo[2.2.1]hept-2-yl)-1,3-dihydro-3,3-ethylenedithio-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one and thence into the title compound.

25 13C Nmr (d₆-acetone) 174.8, 143.2, 141.7, 139.5, 129.6, 128.8, 127.5, 126.6, 125.5, 124.0, 108.6, 63.3, 55.8, 54.0, 53.8, 47.7, 44.0, 39.7, 38.0, 37.4, 36.4, 35.9, 31.0 and 29.3 ppm.

Dihydrochloride, m.p. 253-254°C (dec.).

30

EXAMPLE 101

35 5'-Methyl-1'-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-spiro[1,3-dioxolane-2,3'-[3H]-indol]-2'(1'H)-one
5'-Methyl-spiro[1,3-dioxolane-2,3'-[3H]indol]-2'(1'H)-one (1 equivalent) in dry DMF (5 ml) was added dropwise to sodium hydride (3 equivalents) in dry DMF (2 ml) at 0°C. After 20 minutes, a solution of 4-(2-chloroethyl)-1-(phenylmethyl)piperidine hydrochloride

50

(1.5 equivalents) in dry DMF (15 ml) was slowly added. The mixture was heated to 80°C, stirred at this temperature for 3 hours, and then left at RT overnight. The mixture was evaporated to dryness under reduced pressure and the residue purified by flash chromatography to yield the title compound (53%).
 ^{13}C Nmr (CDCl_3) 173.1, 141.5, 138.5, 132.8, 131.8, 129.2, 128.1, 126.9, 125.6, 124.0, 108.6, 102.1, 65.8, 63.4, 53.6, 37.5, 33.6, 33.4, 32.1 and 20.9 ppm.

10

EXAMPLE 102

5-Methyl-1-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-1H-indole-2,3-dione

The compound of Example 101 (850 mg) in tetrahydrofuran
15 (30 ml) was treated with 3M hydrochloric acid (17 ml).
The mixture was heated under reflux overnight, then
cooled, and neutralised by the addition of aqueous
sodium bicarbonate. The mixture was extracted with
dichloromethane. The extracts were washed, dried and
20 evaporated and the residue was purified by flash
chromatography to give the title compound.

¹³C Nmr (CDCl₃) 183.9, 158.1, 148.6, 138.7, 138.4, 133.5, 129.2, 128.1, 126.9, 125.8, 117.6, 109.9, 63.4, 53.6, 37.9, 33.7, 33.3, 32.1 and 20.7 ppm.

25 Hydrochloride, m.p. 195-197°C.

Following the general methods of Examples 101 and 102 and starting from the appropriately substituted spiro [1,3-dioxolane-2,3'-[3H]indol]-2'(1'H)-one, the compounds of Examples 103 and 104 were prepared.

EXAMPLE 103

5-Methoxy-1-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-1H-indole-2,3-dione

35 ^{13}C Nmr (CDCl_3) 183.7, 158.0, 156.3, 144.5, 138.3, 129.0, 128.0, 126.8, 124.4, 117.9, 110.9, 109.6, 63.2, 55.8, 53.4, 37.8, 33.5, 33.2 and 32.0 ppm.

EXAMPLE 104

5-Cyclohexyl-1-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-1H-indole-2,3-dione

5 ^{13}C Nmr (CDCl₃) 183.8, 158.3, 148.7, 143.9, 136.8,
129.5, 128.2, 127.4, 123.6, 117.7, 109.8, 62.9, 53.2,
43.7, 37.8, 34.2, 33.5, 33.0, 31.4, 26.6 and 25.8 ppm.
Hydrochloride, m.p. 211-213°C.

EXAMPLE 105

10 1,3-Dihydro-5-methyl-1-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-2H-indol-2-one

The compound of Example 102 was reacted according to the general method of Example 99 to give the title compound.

15 ^{13}C Nmr (CDCl₃) 174.6, 142.0, 138.3, 131.4, 129.1,
128.3, 127.9, 126.7, 125.2, 124.6, 107.8, 63.2, 53.5,
37.6, 35.6, 33.7, 33.5, 31.9 and 20.9 ppm.

EXAMPLE 106

20 1,3-Dihydro-5-methoxy-1-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-2H-indol-2-one

The compound of Example 103 was reacted according to the general method of Example 99 to give the title compound.

25 ^{13}C Nmr (CDCl₃) 174.2, 155.5, 138.2, 137.8, 129.0,
128.0, 126.7, 125.9, 112.0, 111.8, 108.2, 63.2, 55.6,
53.4, 37.6, 35.9, 33.7, 33.3 and 32.0 ppm.

EXAMPLE 107

30 5-Cyclohexyl-1,3-dihydro-1-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]-2H-indol-2-one

The compound of Example 104 was reacted according to the general method of Example 99 to give the title compound.

35 ^{13}C Nmr (CDCl₃) 174.8, 142.4, 142.3, 138.4, 129.1, 128.1,
126.8, 125.8, 124.7, 123.1, 107.9, 63.4, 53.6, 44.2,
37.7, 35.9, 34.8, 33.9, 33.5, 32.1, 26.9 and 26.1 ppm.

Following the general method of Example 1 and using the appropriately substituted aniline, the compounds of Examples 108 to 111 were prepared.

5

EXAMPLE 108**5-Cyclopentyl-1H-indole-2,3-dione**

M.p. 138-140°C.

 ^{13}C Nmr (CDCl_3) 183.8, 160.2, 147.5, 142.7, 137.8, 124.0, 117.9, 112.5, 45.1, 34.4 and 25.3 ppm.

10

EXAMPLE 109**7-Cyclopentyl-1H-indole-2,3-dione** ^1H Nmr (CDCl_3) 1.5 - 1.9 (6H, m), 2.1 (2H, m), 3.0 (1H, m), 7.05 (1H, t), 7.45 (2H, m) and 9.0 (1H, br s) ppm.

15

EXAMPLE 110**5-Cycloheptyl-1H-indole-2,3-dione** ^1H Nmr (CDCl_3) 1.5 - 1.9 (12H, m), 2.65 (1H, m), 6.85 (1H, d), 7.4 (1H, dd), 7.45 (1H, d) and 8.6 (1H, br s) ppm.

20

EXAMPLE 111**7-Cycloheptyl-1H-indole-2,3-dione** ^1H Nmr (CDCl_3) 1.5 - 2.0 (12H, m), 2.65 (1H, m), 7.05 (1H, t), 7.45 (2H, d) and 8.6 (1H, br s) ppm.

30

Following the general method of Example 3 and using the appropriately substituted 1H-indole-2,3-dione, the compounds of Examples 112 to 114 were prepared.

EXAMPLE 112**5-Cyclopentyl-1,3-dihydro-2H-indol-2-one** ^1H Nmr (CDCl_3) 1.5 - 1.9 (6H, m), 2.05 (2H, m), 2.95 (1H, m), 3.55 (2H, s), 6.8 (1H, d), 7.1 (2H, m) and 8.6 (1H, br s) ppm.

EXAMPLE 113**5-Cycloheptyl-1,3-dihydro-2H-indol-2-one**

5 ¹H Nmr (CDCl₃) 1.5 - 2.0 (12H, m), 2.65 (1H, m), 3.55
ppm.

EXAMPLE 114**7-Cycloheptyl-1,3-dihydro-2H-indol-2-one**

10 ¹H Nmr (CDCl₃) 1.5 - 2.0 (12H, m), 2.65 (1H, m), 3.55
(2H, s), 6.95 - 7.1 (3H, m) and 8.4 (1H, br s) ppm.

EXAMPLE 115**1,3-Dihydro-5-(1-pyrrolidinyl)-2H-indol-2-one**

15 2-Methyl-4-(1-pyrrolidinyl)-aniline was converted into
the N-(tert-butoxycarbonyl) derivative and thence into
the title compound using the methodology of R.D. Clark
et al, *Synthesis* 1991, 871-878.

20 ¹³C Nmr (d₆-DMSO) 175.7, 143.8, 133.1, 126.7, 110.0,
109.4, 109.1, 47.8, 36.2 and 24.7 ppm.

20

EXAMPLE 116**1,3-Dihydro-5-(1-piperidinyl)-2H-indol-2-one**

The title compound was prepared from 2-methyl-4-(1-piperidinyl)-aniline using the method of Example 115.

25 M.p. 154-156°C.

¹³C Nmr (CDCl₃) 177.8, 148.3, 136.2, 126.2, 117.0,
115.4, 109.9, 52.6, 36.7, 25.8 and 24.0 ppm.

EXAMPLE 117**5-Diethylamino-1,3-dihydro-2H-indol-2-one**

The title compound was prepared from 4-diethylamino-2-methylaniline using the method of Example 115.

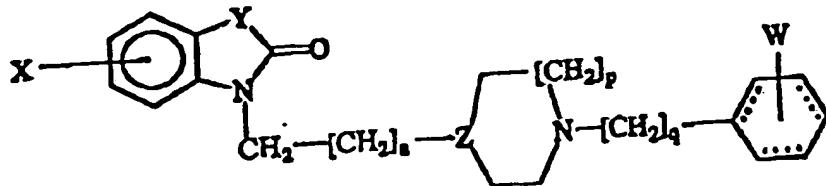
M.p. 122-124°C.

30 ¹H Nmr (CDCl₃) 1.1 (6H, t), 3.25 (4H, q), 3.55 (2H, s),
6.6 (1H, dd), 6.75 (2H, m) and 9.0 (1H, br s) ppm.

CLAIMS

1. A compound having the general formula (1)

5



(1)

10

wherein:

n is 1, 2 or 3;

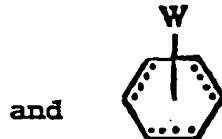
15 p is 1 or 2;

q is 1 or 2;

X represents one or more substituents independently selected from hydrogen, lower alkyl, aryl, aryloxy, CN, lower alkoxy, halogen, hydroxy, nitro, trifluoromethyl, alkylsulphonamido, NHCOR where R is lower alkyl or aryl, NR₁R₂ where R₁ and R₂ are independently hydrogen or lower alkyl or together form a ring, CO₂R where R is lower alkyl, or cycloalkyl, cycloalkenyl or bicycloalkyl either optionally further substituted by lower alkyl;

30 Y is CO or CR₃R₄ where R₃ and R₄ are independently hydrogen, lower alkyl, lower alkoxy or together form a cyclic acetal;

35 Z is N or CH;



and represents an optically substituted phenyl or cyclohexyl group; wherein

W r presents one or more substituents independently selected from hydrogen, lower alkyl, lower alkoxy or halogen;

5

stereo and optical isomers and racemates thereof where such isomers exist, as well as pharmaceutically acceptable acid addition salts thereof and solvates thereof;

10

with the provisos that the compound wherein n=1, p=1,

15

q=1, X=H, Y=CO, Z=N and



= unsubstituted

phenyl and the compound wherein n=2, p=1, q=1, X=H,

20

Y=CO, Z=N and

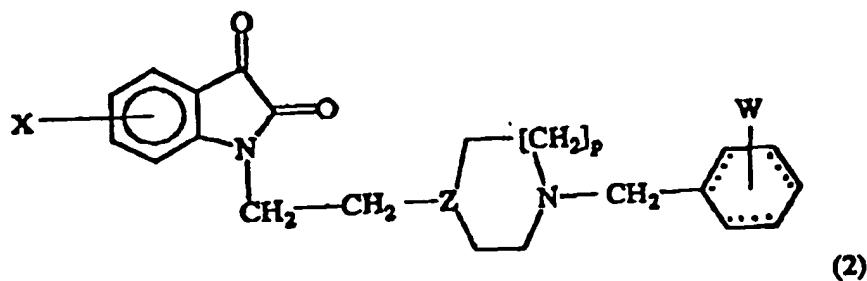


= 4-chlorophenyl are excluded.

2. A compound according to claim 1 having the general formula (2)

25

30

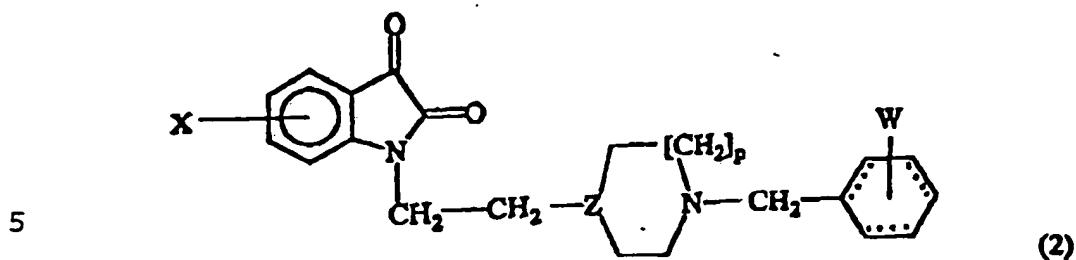


wherein Z=N and p, X and W are as defined in claim 1.

35

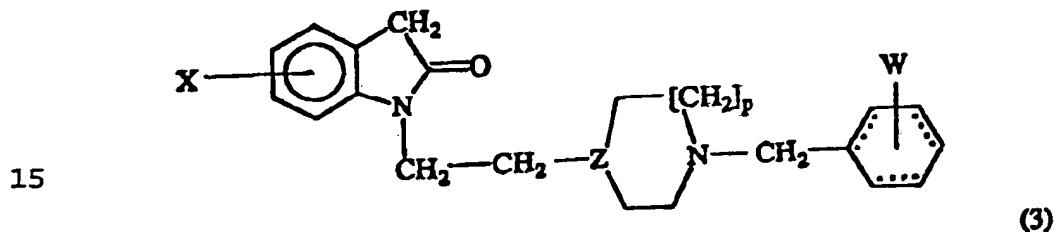
3. A compound according to claim 1 having the general formula (2)

56



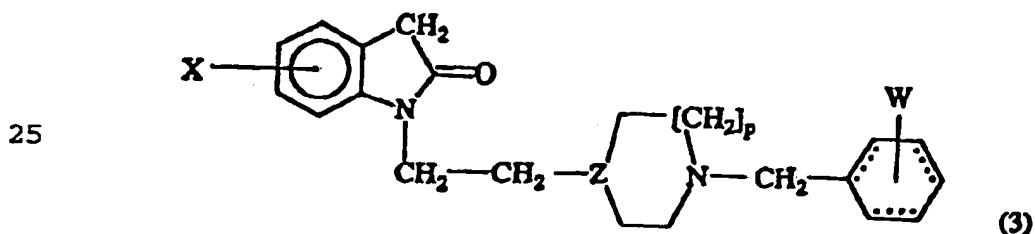
wherein $Z=CH$ and p , X and W are as defined in claim 1.

- 10 4. A compound according to claim 1 having the general formula (3)



wherein $Z=N$ and p , X and W are as defined in claim 1.

- 20 5. A compound according to claim 1 having the general formula (3)



wherein $Z=CH$ and p , X and W are as defined in claim 1.

- 30 6. A compound according to any one of claims 1-5
wherein the X substituent is at the 5-position.

- 35 7. A compound according to any one of claims 2 - 6
wherein
 p is 1,
 W is hydrogen or F, and

X is lower alkyl, low r alkoxy, cycloalkyl, F, aryl, or NR₁R₂ where R₁ and R₂ are independently hydrogen or lower alkyl or together form a ring.

5 8. A compound according to claim 7 wherein
W is H or 4-F, and

X is methyl, ethyl, methoxy, ethoxy, C₅ to C₇
cycloalkyl, F, aryl, especially phenyl, or NR₁R₂,
especially 1-pyrrolidinyl or 1-piperidinyl.

10

9. A compound according to claim 1 consisting of

- 1,3-Dihydro-5-methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one,
- 5-Cyclohexyl-1,3-dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one,

15

- 1,3-Dihydro-1-[2-[4-[(4-fluorophenyl)methyl]-1-piperazinyl]ethyl]-5-methyl-2H-indol-2-one,
- 5-Cyclohexyl-1,3-dihydro-1-[2-[4-[4-fluoro-phenyl)methyl]-1-piperazinyl]ethyl]-2H-indol-2-one,

20

- 5-Methyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione,
- 1-[2-[4-[(4-Fluorophenyl)methyl]-1-piperazinyl]ethyl]-5-methyl-1H-indole-2,3-dione,
- 5-Cyclohexyl-1-[2-[4-(phenylmethyl)-1-

25

- piperazinyl]ethyl]-1H-indole-2,3-dione,
- 5-Fluoro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-1H-indole-2,3-dione,
- 1,3-Dihydro-5-fluoro-1-[2-[4-(phenylmethyl)-1-

30

- piperazinyl]ethyl]-2H-indol-2-one,
- 1,3-Dihydro-5-phenyl-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-2H-indol-2-one,
- 1,3-Dihydro-1-[2-[4-(phenylmethyl)-1-piperazinyl]ethyl]-5-(1-piperidinyl)-2H-indol-2-one,
- 5-Cyclohexyl-1,3-dihydro-1-[2-[1-(phenylmethyl)-4-

35

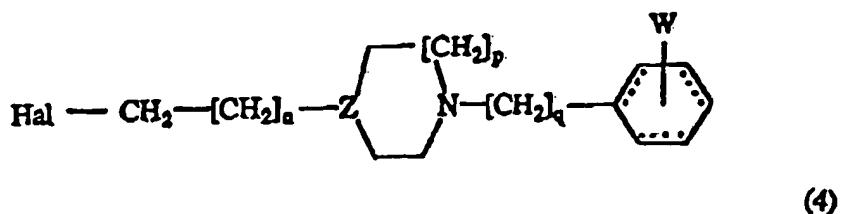
- piperidinyl]ethyl]-2H-indol-2-one

and pharmaceutically acceptabl acid addition salts r

solvates the reof.

10. A process for preparing a compound according to
any one of the preceding claims by
5 (a) reacting a compound of the general formula (4)

10



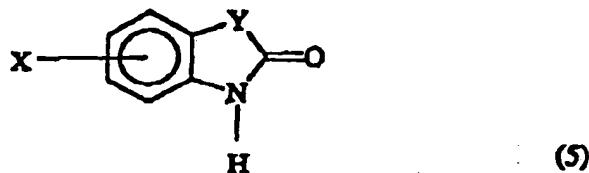
(4)

wherein Z , W , n , p and q are as defined in claim 1 and
Hal is halogen,

15

with a compound of the general formula (5)

20



(5)

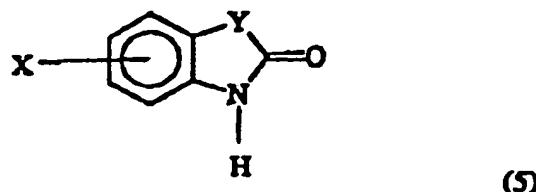
wherein X and Y are as defined in claim 1,

25

or, in the case where $Z=N$, by

(b) treating a compound of the general formula (5)

30



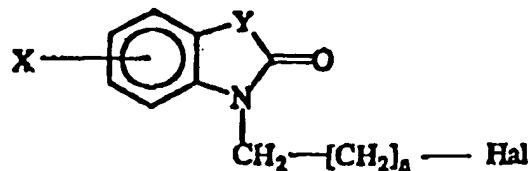
(5)

wherein X and Y are as defined in claim 1,

35

with a $1,(n+1)$ -dihaloalkane to obtain a compound of the
general formula (6)

5

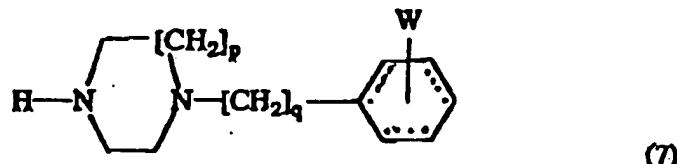


wherein X, Y and n are as defined in claim 1 and Hal is halogen,

10

and reacting the compound of the general formula (6) with a compound of the general formula (7)

15

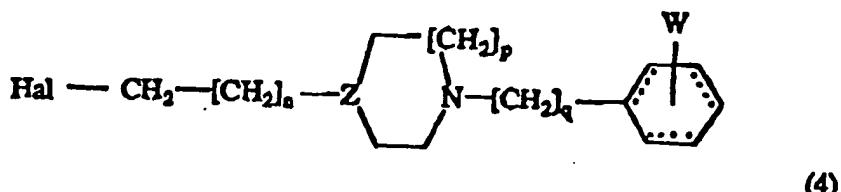


wherein W, p and q are as defined in claim 1.

20

11. A compound of the general formula (4)

25



wherein Z is N or CH, Hal is halogen n=p=q=1 and W=Me, OMe or F

30

or

=cyclohexyl, with

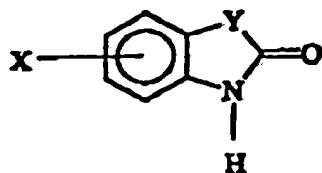
the proviso that the compound where Z=N and = 2-methylphenyl is excluded.

35

12. A compound of the general formula (5)



60



5 wherein

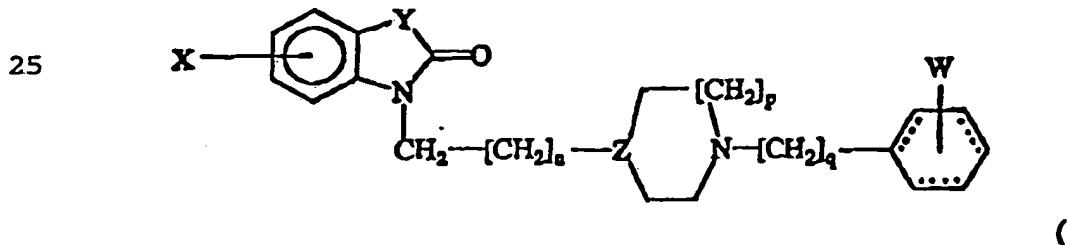
X is cycloalkyl, cycloalkenyl or bicycloalkyl, either
optionally further substituted by lower alkyl or X is
10 $-N\left(\text{CH}_2\right)_n$ where n= 4 to 7

15 and Y is CH_2 or CO or $\text{C}\left(\text{CH}_2\right)_m$ where m= 2 to 4,

with the proviso that the compound where X=5-cyclohexyl
and Y=CO is excluded.

20

13. A pharmaceutical formulation containing a compound
having the general formula (1)



30 wherein:

n is 1, 2 or 3;

p is 1 or 2;

35

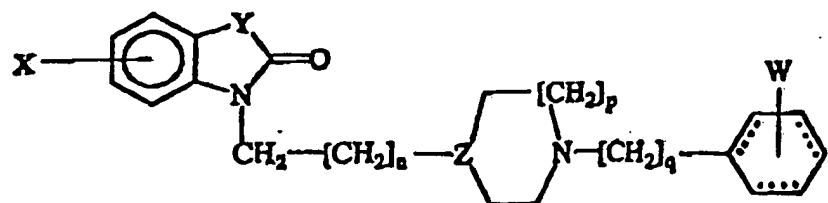
q is 1 or 2;

40 X represents one or more substituents independently selected from hydrogen, lower alkyl, aryl, aryloxy, CN, lower alkoxy, halogen, hydroxy, nitro, trifluoromethyl,

- alkylsulphonamido,
 NHCOR where R is lower alkyl or aryl,
 NR_1R_2 where R_1 and R_2 are independently hydrogen or
 lower alkyl or together form a ring,
 5 CO_2R where R is lower alkyl,
 or cycloalkyl, cycloalkenyl or bicycloalkyl either
 optionally further substituted by lower alkyl;
- Y is CO or CR_3R_4 where R_3 and R_4 are independently
 10 hydrogen, lower alkyl, lower alkoxy or together form a
 cyclic acetal;
- Z is N or CH;
 15 
 and  represents an optionally substituted
 phenyl or cyclohexyl group; wherein
- W represents one or more substituents independently
 20 selected from hydrogen, lower alkyl, lower alkoxy or
 halogen;
- 25 stereo and optical isomers and racemates thereof where
 such isomers exist, as well as pharmaceutically
 acceptable acid addition salts thereof and solvates
 thereof;
- 30 with the proviso that the compound wherein $n=2$, $p=1$,
 $q=1$, $\text{X}=\text{H}$,
- 35 $\text{Y}=\text{CO}$, $\text{Z}=\text{N}$ and  = 4-chlorophenyl is excluded,
 as active ingredient and a pharmaceutically acceptable
 carrier.

14. A compound having the general formula (1)

5



10

wherein:

n is 1, 2 or 3;

15 p is 1 or 2;

q is 1 or 2;

20 X represents one or more substituents independently selected from hydrogen, lower alkyl, aryl, aryloxy, CN, lower alkoxy, halogen, hydroxy, nitro, trifluoromethyl, alkylsulphonamido,
 NHCOR where R is lower alkyl or aryl,
 NR₁R₂ where R₁ and R₂ are independently hydrogen or
 25 lower alkyl or together form a ring,
 CO₂R where R is lower alkyl,
 or cycloalkyl, cycloalkenyl or bicycloalkyl either optionally further substituted by lower alkyl;

30 Y is CO or CR₃R₄ where R₃ and R₄ are independently hydrogen, lower alkyl, lower alkoxy or together form a cyclic acetal;

Z is N or CH;

35

and

represents an optionally substituted

phenyl or cyclohexyl group; wherein

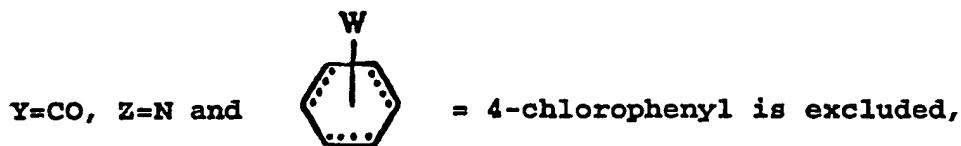
W represents one or more substituents independently selected from hydrogen, lower alkyl, lower alkoxy or halogen;

5 halogen;

stereo and optical isomers and racemates thereof where such isomers exist, as well as pharmaceutically acceptable acid addition salts thereof and solvates thereof;

with the proviso that the compound wherein $n=2$, $p=1$,
 $q=1$, $X=H$,

15



20

for use in therapy.

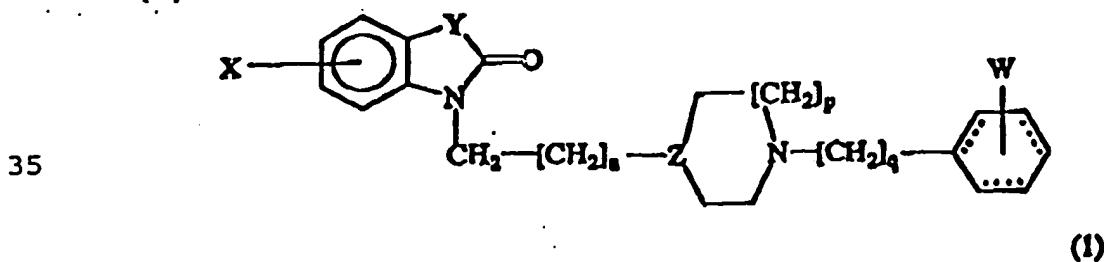
15. A compound as defined in claim 14 for use as an agent for the treatment of conditions which involve a decreased cholinergic function.

25

16. A compound as defined in claim 14 for use as an agent for prevention or treatment of cognitive dysfunctions.

30

17. The use of a compound having the general formula
(1)



wherein:

5 n is 1, 2 or 3;

p is 1 or 2;

q is 1 or 2;

10

X represents one or more substituents independently selected from hydrogen, lower alkyl, aryl, aryloxy, CN, lower alkoxy, halogen, hydroxy, nitro, trifluoromethyl, alkylsulphonamido,

15 NHCOR where R is lower alkyl or aryl,

NR₁R₂ where R₁ and R₂ are independently hydrogen or lower alkyl or together form a ring,

CO₂R where R is lower alkyl,

or cycloalkyl, cycloalkenyl or bicycloalkyl either optionally further substituted by lower alkyl;

20

Y is CO or CR₃R₄ where R₃ and R₄ are independently hydrogen, lower alkyl, lower alkoxy or together form a cyclic acetal;

25

Z is N or CH;



30

and represents an optionally substituted

phenyl or cyclohexyl group; wherein

35

W represents one or more substituents independently selected from hydrogen, lower alkyl, lower alkoxy or halogen;

stereo and optical isomers and racemates thereof where

such isomers exist, as well as pharmaceutically acceptable acid addition salts thereof and solvates thereof for the manufacture of a medicament for the treatment of conditions which involve a decreased cholinergic function.

5 18. The use according to claim 17 for the treatment of conditions such as glaucoma or myasthenia gravis.

10 19. The use according to claim 17 for the manufacture of a medicament for the prevention or treatment of cognitive dysfunctions.

15 20. The use according to claim 19 for the prevention or treatment of cognitive dysfunctions associated with ageing.

20 21. The use according to claim 19 for the prevention or treatment of cognitive dysfunctions associated with conditions such as Alzheimer's Disease, Senile and related Dementias, Parkinson's Disease, Down's Syndrome and Huntington's Chorea.

25 22. A method for the prevention or treatment of decreased cholinergic function by administering to a host in need of such a treatment a sufficient amount of a compound according to claim 1.

30 23. A method for the prevention or treatment of cognitive dysfunctions by administering to a host in need of such a treatment a sufficient amount of a compound according to claim 1.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 92/00873

A. CLASSIFICATION OF SUBJECT MATTER

IPC5: C07D 209/34, C07D 209/38, C07D 295/067, C07D 491/113, C07D 403/00,
A61K 31/495

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC5: C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP, A1, 0010398 (FUJISAWA PHARMACEUTICAL CO. LTD.), 30 April 1980 (30.04.80) --	1-11,13-19
A	Chemical Abstracts, volume 98, no. 3, 17 January 1983, (Columbus, Ohio, US), F. Collino et al: "Mannich bases of benzimidazoles, benzotriazoles and other analogous compounds, with pharmacological activity", see pages 508, abstract 16650w, & Boll. Chim. Farm. 1982, 121(5), 221-29, see reg. nr 83991-57-5 --	1-11,13-19
A	US, A, 4895841 (SUGIMOTO ET AL.), 23 January 1990 (23.01.90), see columns 1-12, 57-60 and example 10 --	1-11,13-19

 Further documents are listed in the continuation of Box C. See patent family annex.

- * Special categories of cited documents:
- "A" document defining the general state of the art which is not considered to be of particular relevance
- "B" earlier document but published on or after the international filing date
- "I" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed
- "T" later document published after the international filing date or priority date and not to conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
11 May 1993	14 -05- 1993
Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Facsimile No. + 46 8 666 02 86	Authorized officer Göran Karlsson Telephone No. + 46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 92/00873

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO, A1, 9102725 (SYNTHESES ET RECHERCHES), 7 March 1991 (07.03.91), see example 7	1-11,13-19
A	New England journal of medicine, Volume 315, No 20, November 1986, William Koopmans Summers, M.D. et al, "Oral tetrahydroaminoacridine in long-term treatment of senile dementia, Alzheimer type" page 1241 - page 1245	1-11,13-19

INTERNATIONAL SEARCH REPORT

I: National application No.

PCT/SE 92/00873

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: 20-23
because they relate to subject matter not required to be searched by this Authority, namely:
A method for treatment of the human or animal body by therapy,
see rule 39.1.
2. Claims Nos.: 1-7, 10, 13-19
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
Claims 1-7, 10 and 13-19 are too broadly formulated to permit a meaningful search, see Article 6. The search has thus been limited to the compounds considered to be most relevant.
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

See annexed sheet !

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.: 12
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
- No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT
Information on patent family members

31/03/93

International application No.
PCT/SE 92/00873

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP-A1- 0010398	30/04/80	SE-T3- 0010398 AU-A- 5157679 US-A- 4382934		17/04/80 10/05/83
US-A- 4895841	23/01/90	AU-B- 627151 EP-A- 0296560		20/08/92 28/12/88
WO-A1- 9102725	07/03/91	CA-A- 2064999 EP-A- 0487623		26/02/91 03/06/92

THIS PAGE BLANK (USPTO)